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**Scheller et al.**

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(54) **CANNULA INGRESS SYSTEM**

2017/3482; A61B 2217/005; A61B 2217/007; A61B 17/3462; A61F 9/007; A61M 2205/582; A61M 39/22; A61M 3/0283

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See application file for complete search history.

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(73) Assignee: **Katalyst Surgical, LLC**, Chesterfield, MO (US)

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(65) **Prior Publication Data**

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**Related U.S. Application Data**

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(60) Provisional application No. 62/062,678, filed on Oct. 10, 2014.

(57) **ABSTRACT**

(51) **Int. Cl.**

<b>A61M 3/02</b>	(2006.01)
<b>A61F 9/007</b>	(2006.01)
<b>A61M 39/22</b>	(2006.01)
<b>A61B 17/34</b>	(2006.01)

A cannula ingress system may include a tip stabilization mechanism having a tip stabilization mechanism distal end and a tip stabilization mechanism proximal end, a fixation mechanism, a hypodermic tube having a hypodermic tube distal end and a hypodermic tube proximal end, and a tip having a tip distal end and a tip proximal end. The tip may be disposed within the hypodermic tube wherein the tip distal end extends from the hypodermic tube distal end. The fixation mechanism may be disposed within a fixation mechanism channel of the tip stabilization mechanism. The tip stabilization mechanism may be disposed over the tip and the hypodermic tube distal end.

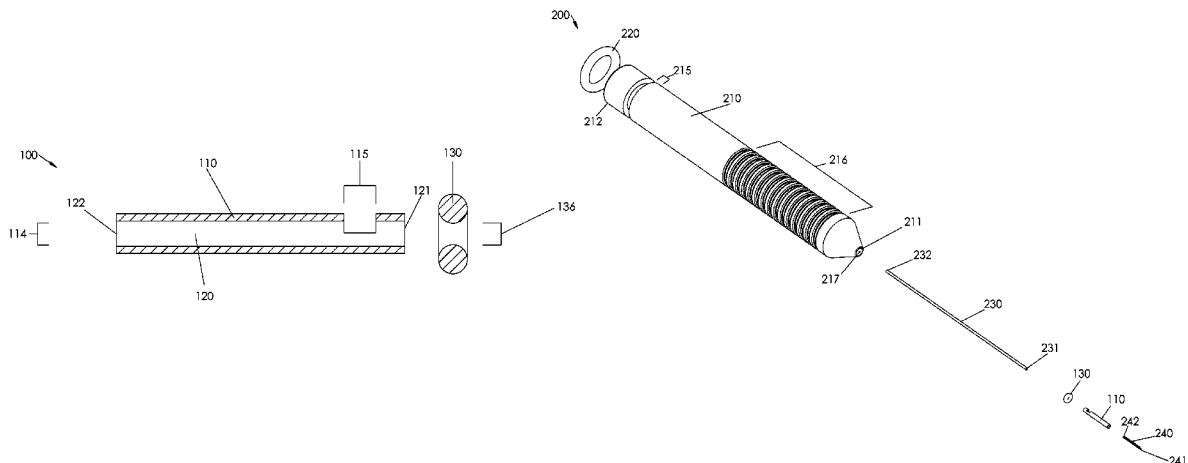
(52) **U.S. Cl.**

CPC ..... **A61M 3/0283** (2013.01); **A61F 9/007** (2013.01); **A61M 39/22** (2013.01); **A61B 17/3421** (2013.01); **A61B 17/3462** (2013.01); **A61M 2205/582** (2013.01)

(58) **Field of Classification Search**

CPC ..... A61B 17/3421; A61B 17/3498; A61B

**20 Claims, 29 Drawing Sheets**



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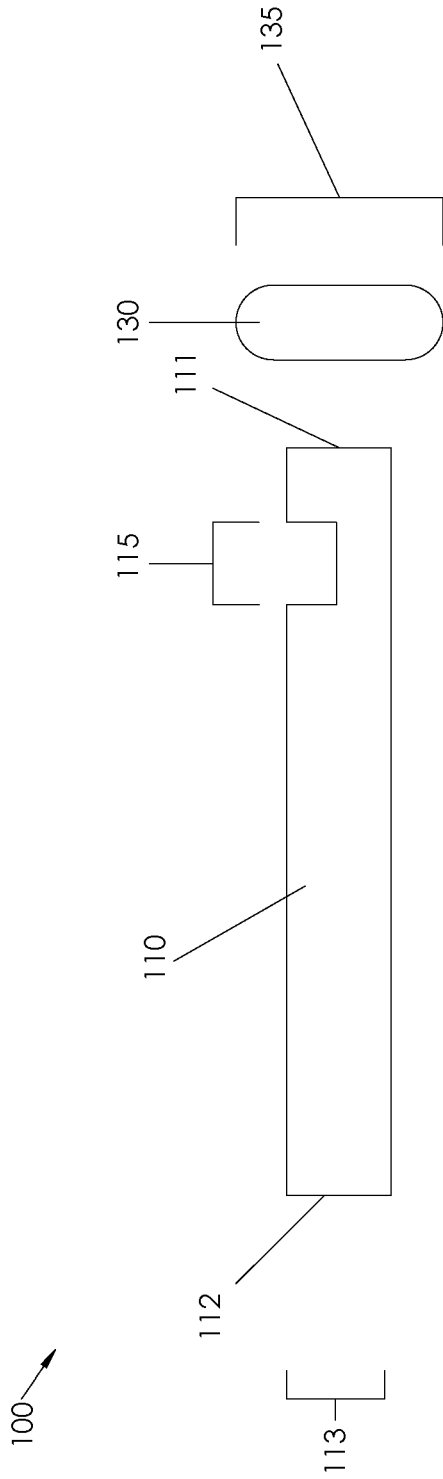


FIG. 1A

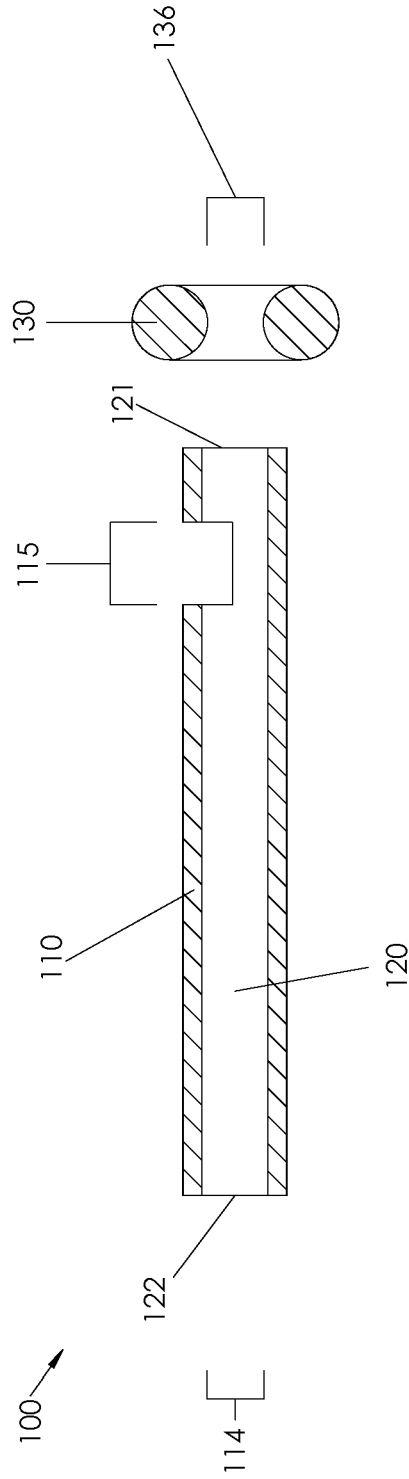


FIG. 1B

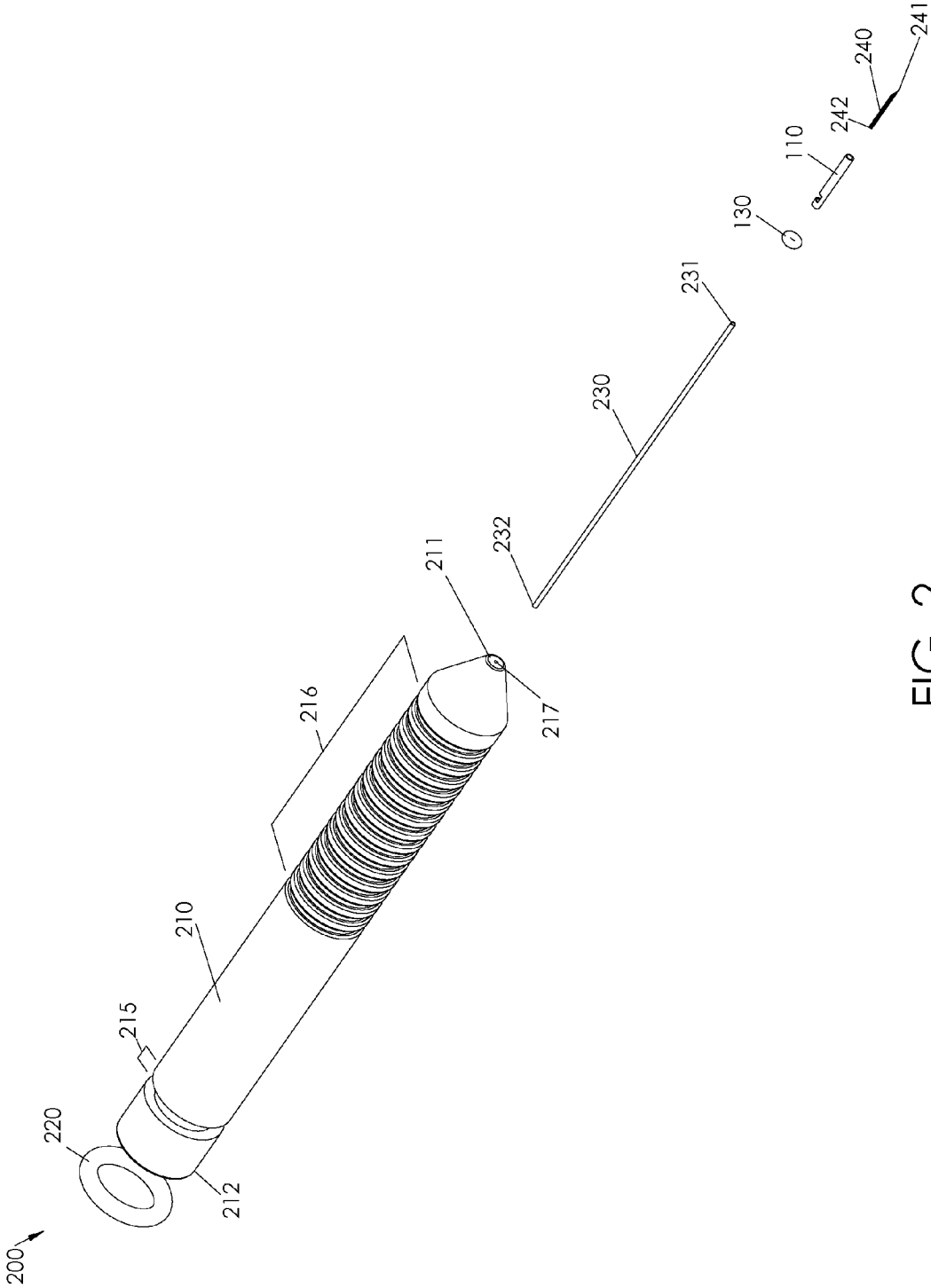


FIG. 2

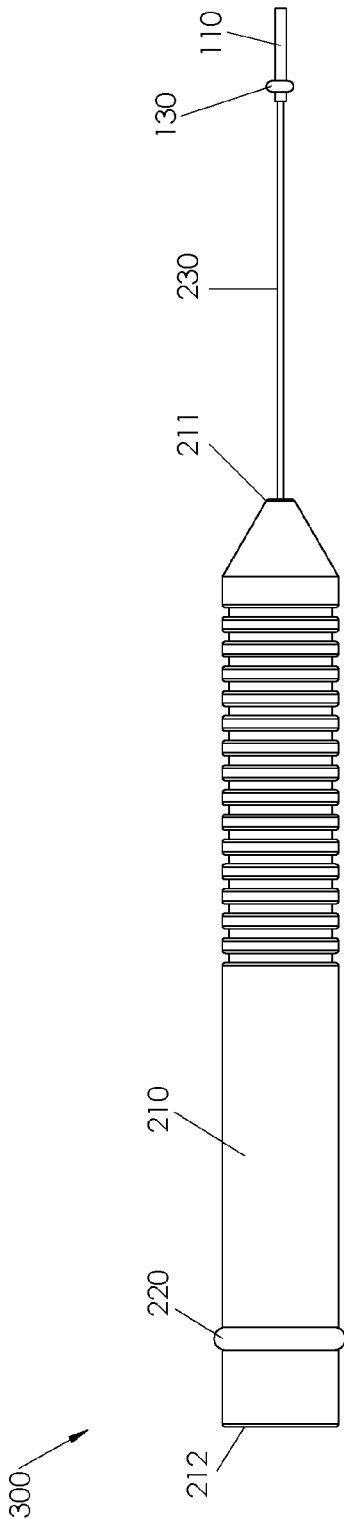


FIG. 3A

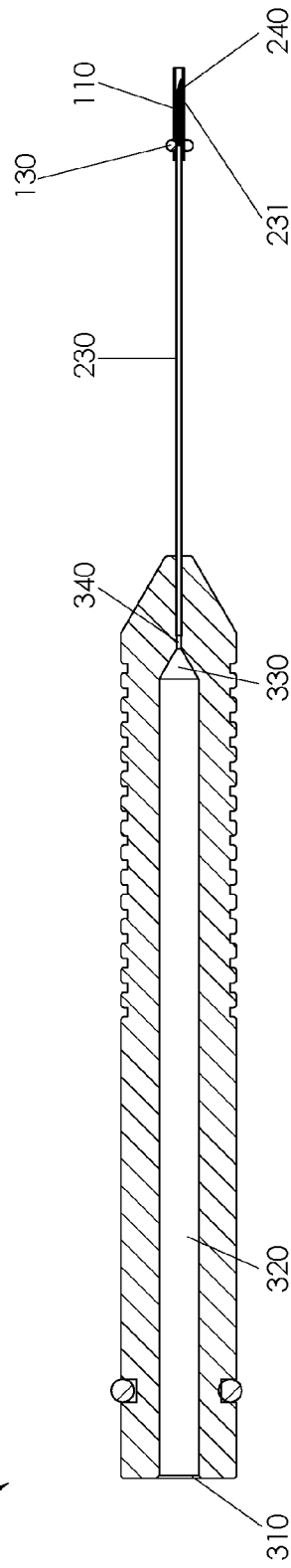


FIG. 3B

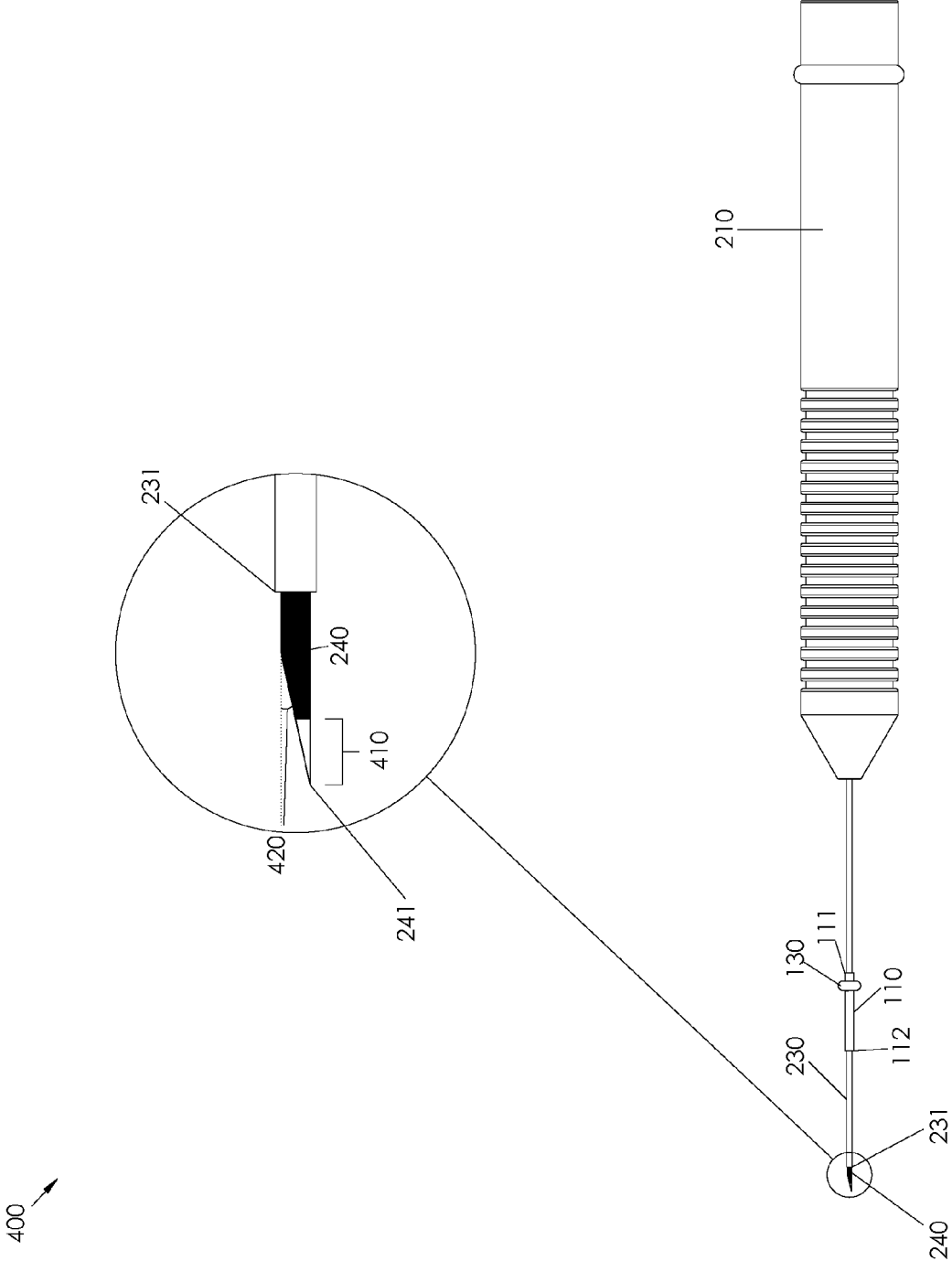


FIG. 4

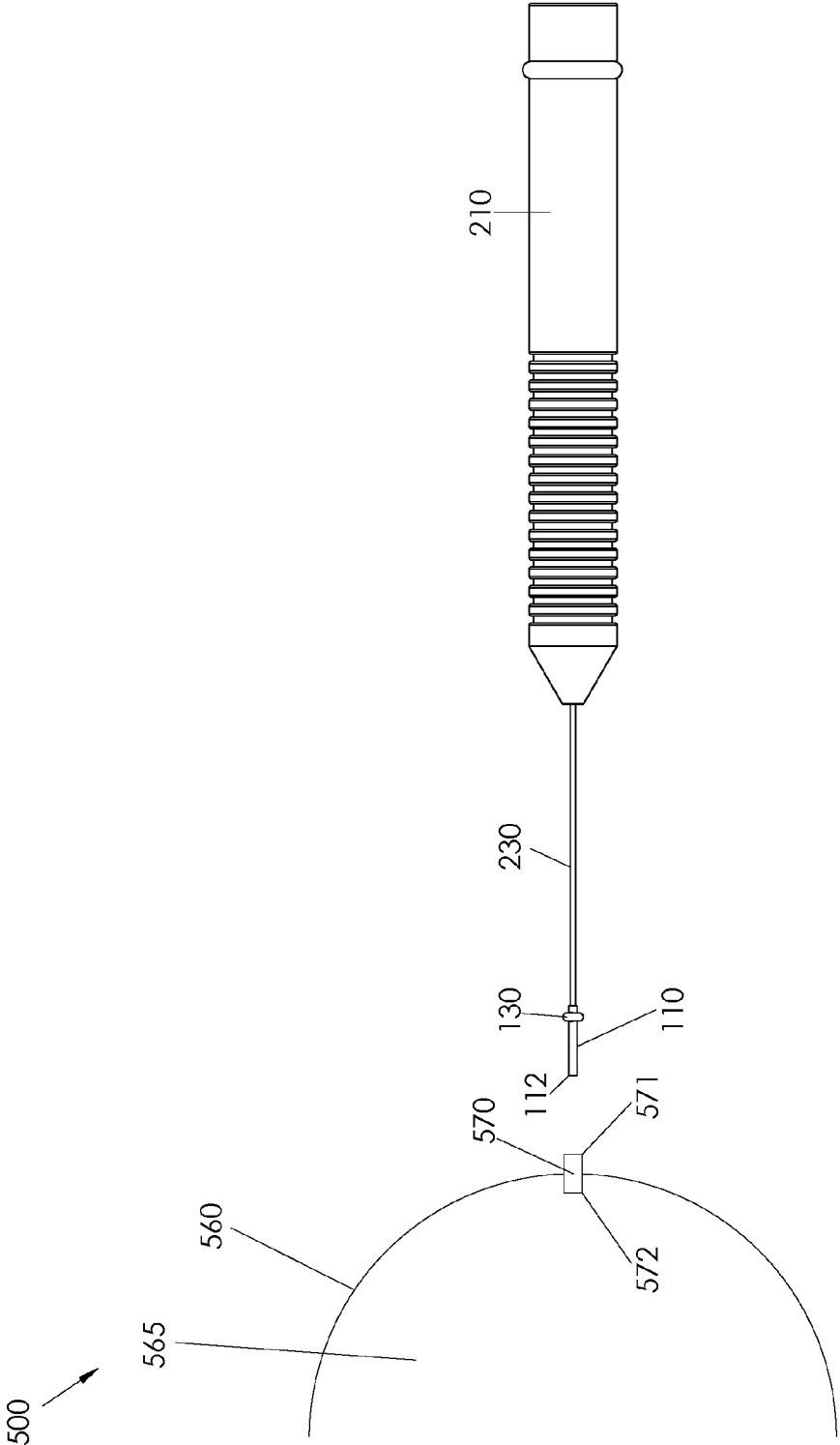


FIG. 5A

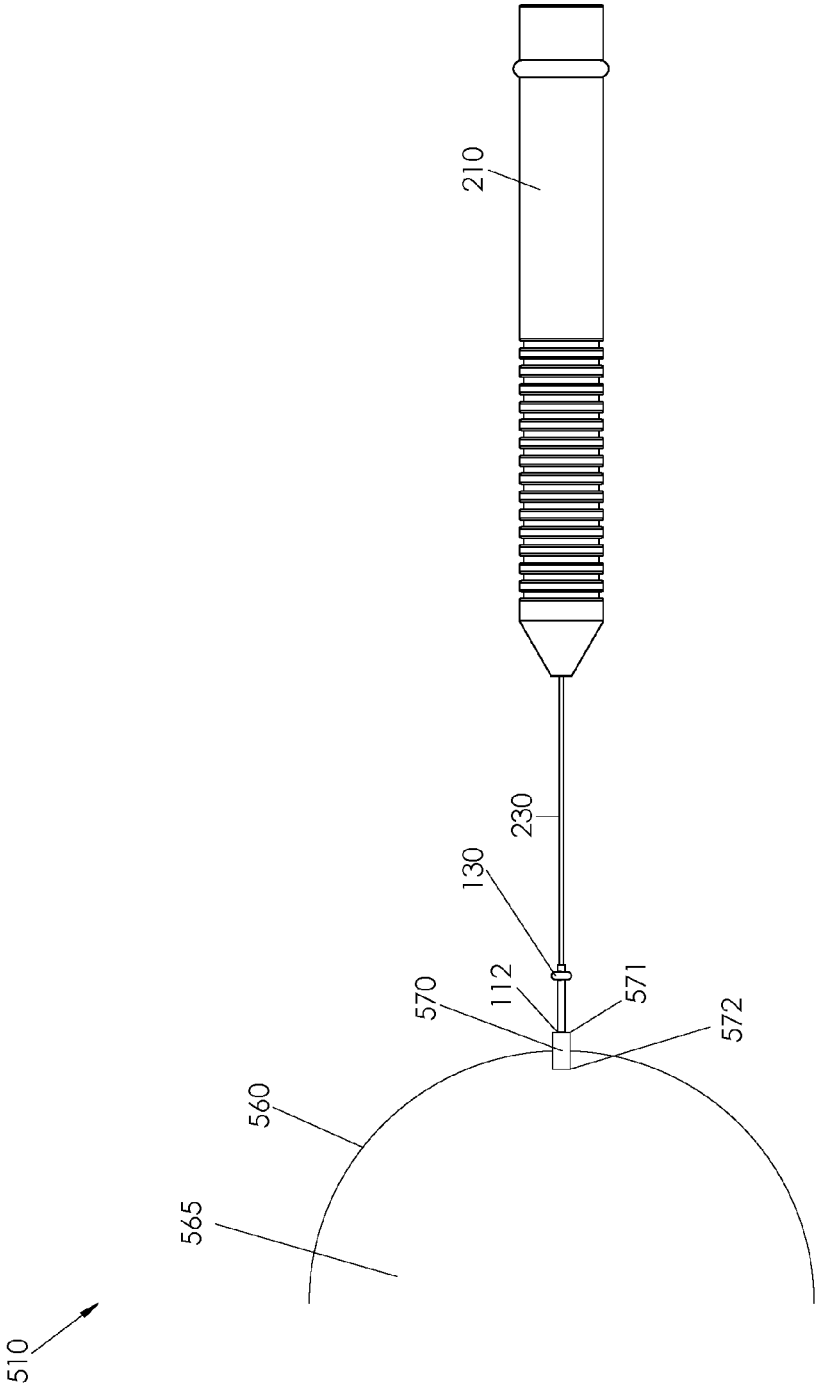


FIG. 5B

520 ↗

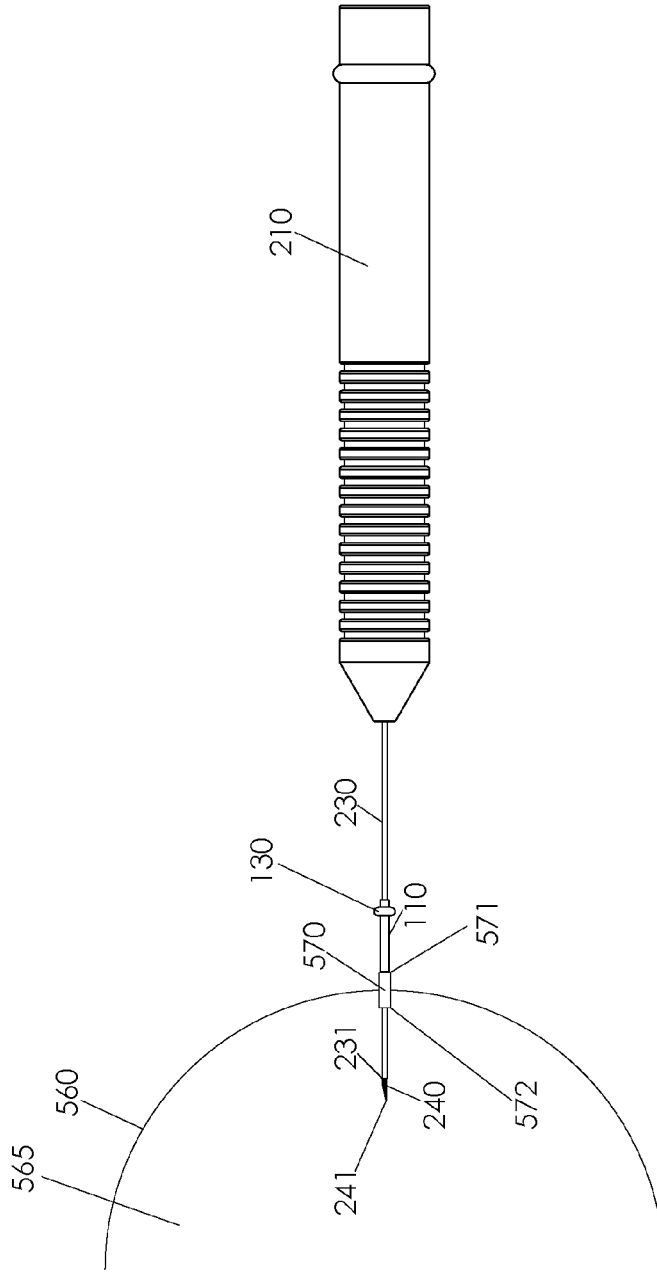


FIG. 5C

530

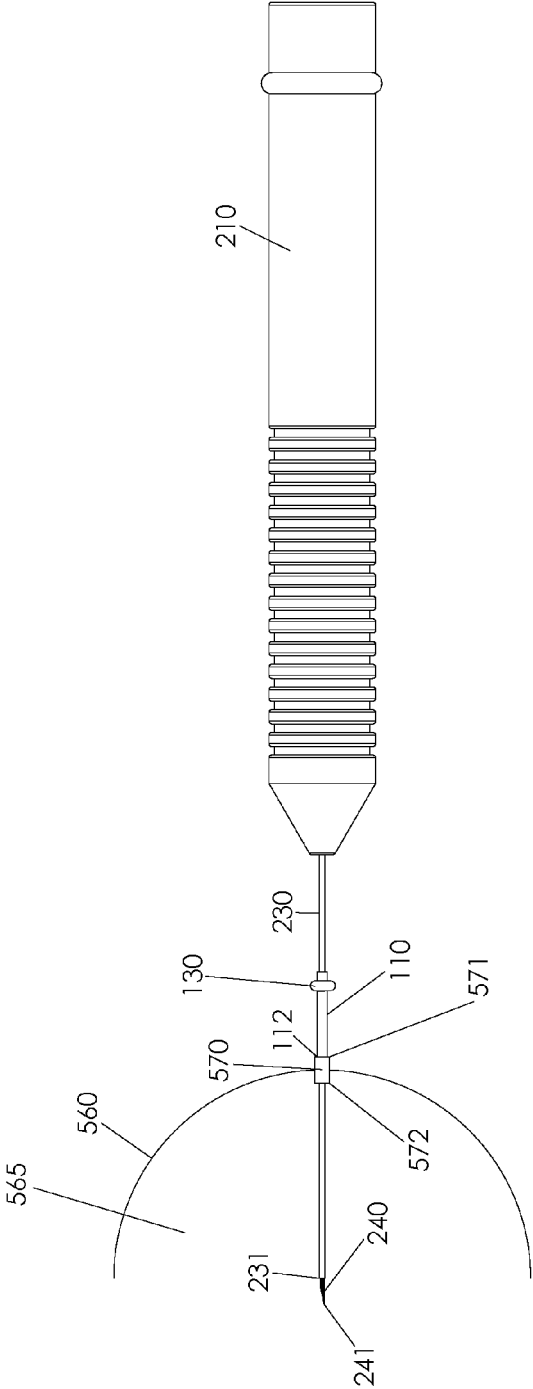


FIG. 5D

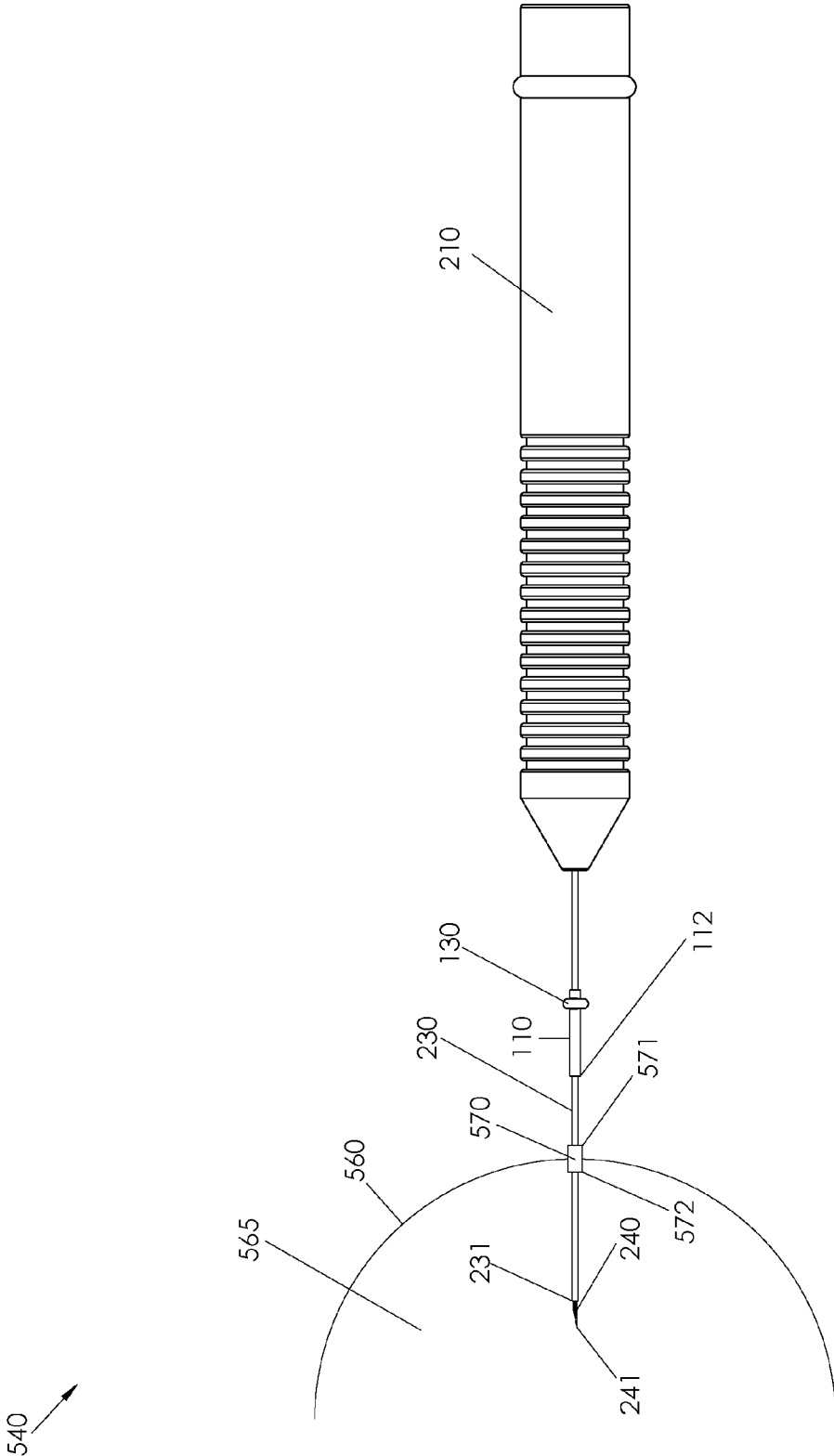


FIG. 5E

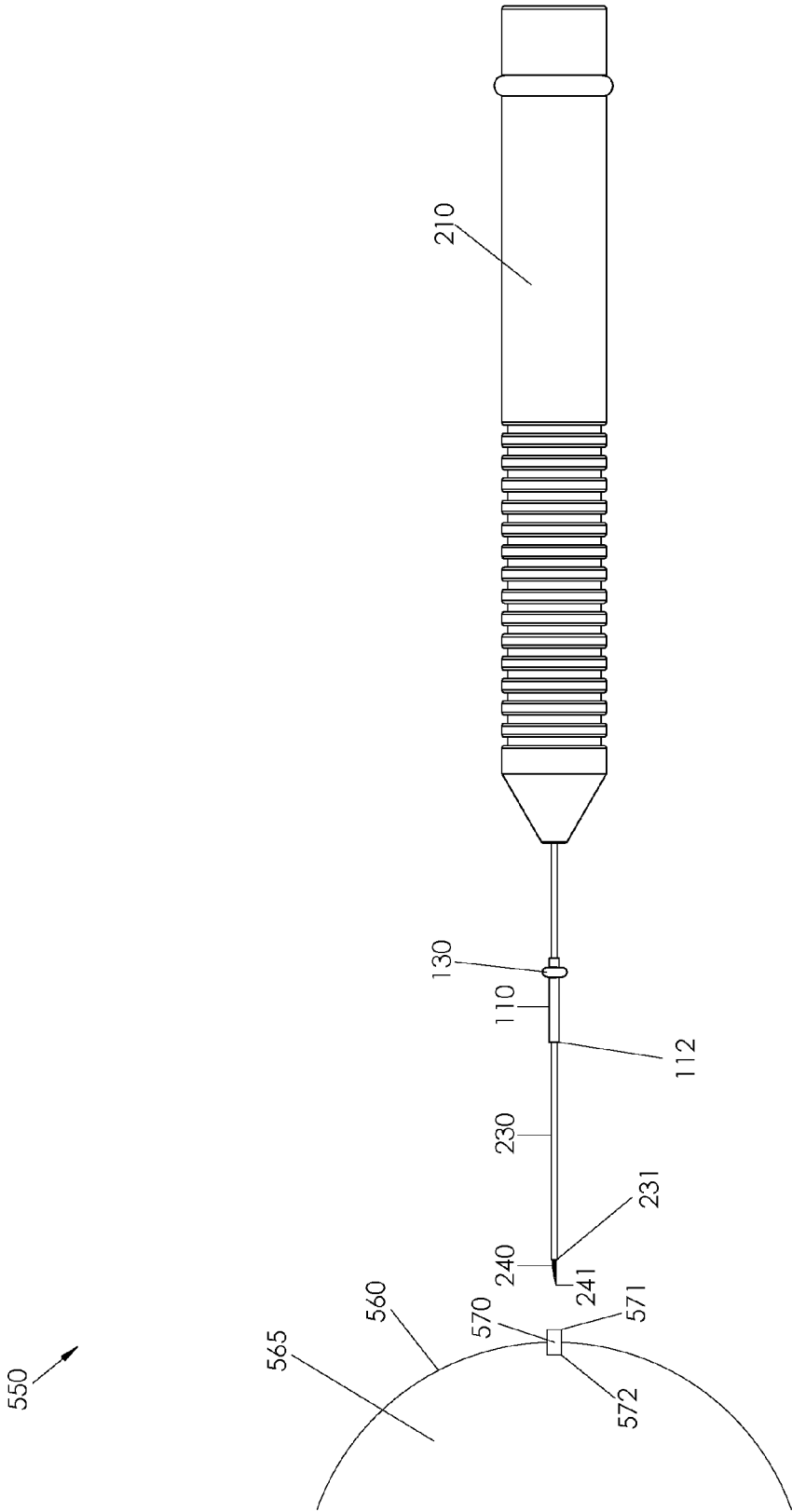


FIG. 5F

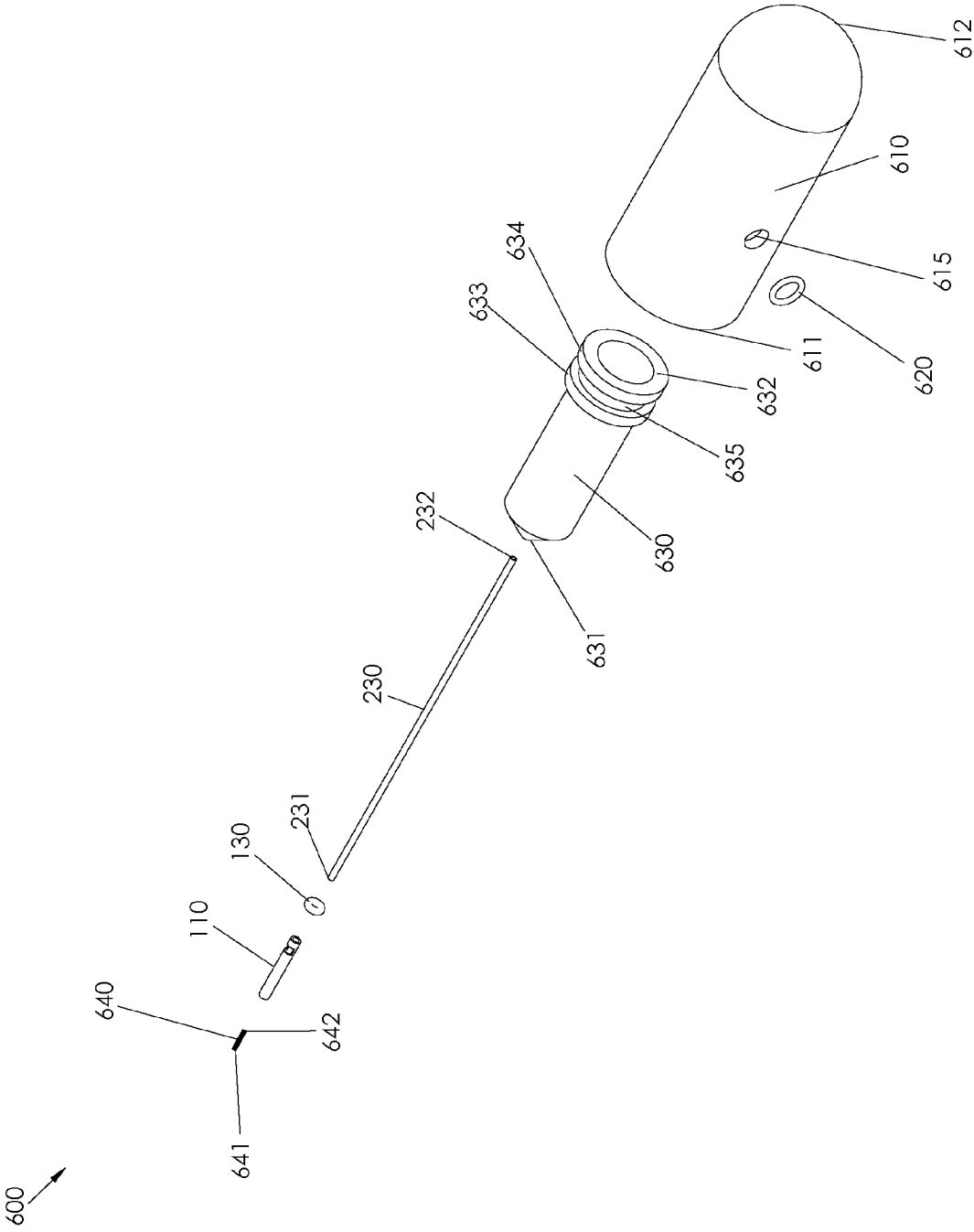


FIG. 6

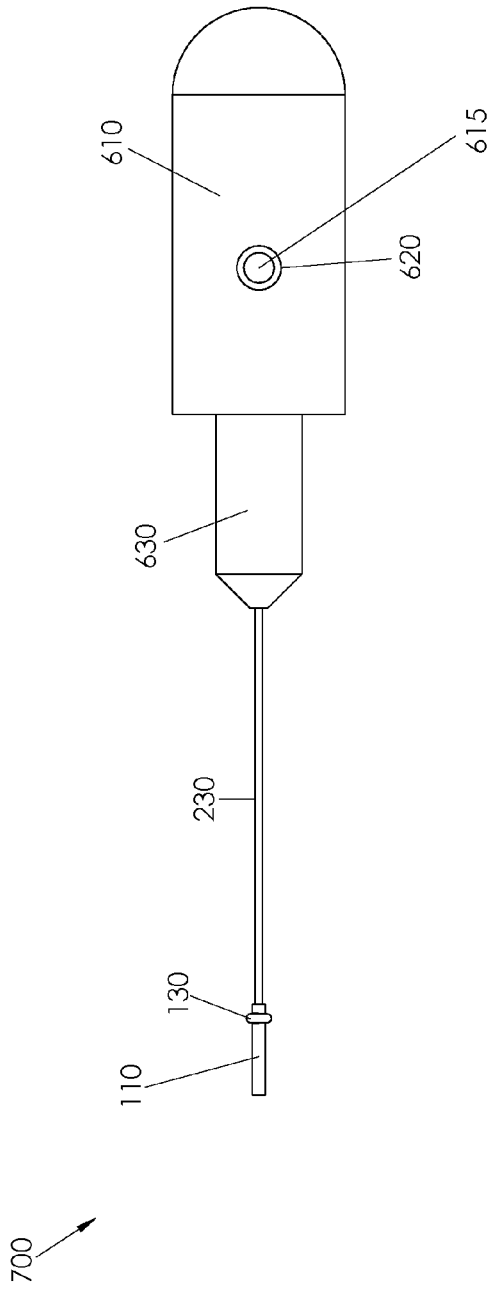


FIG. 7A

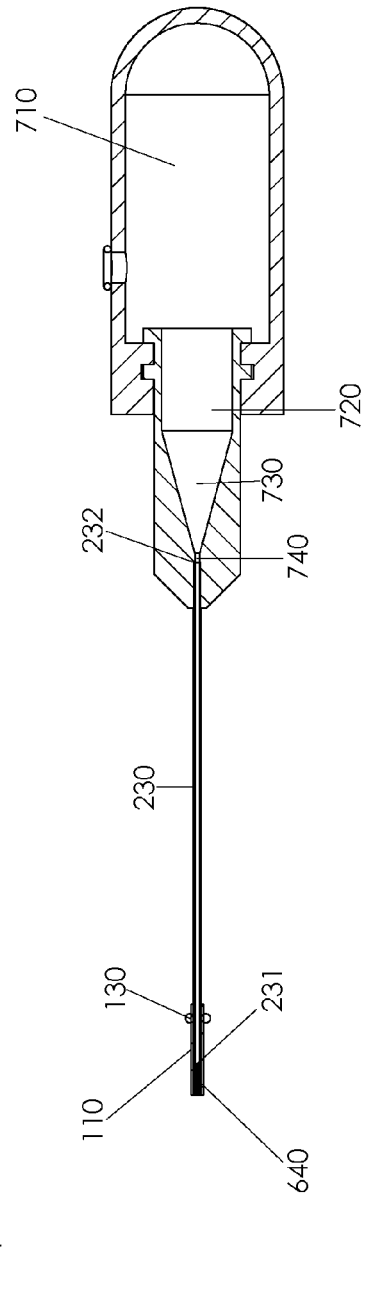


FIG. 7B

800 ↗

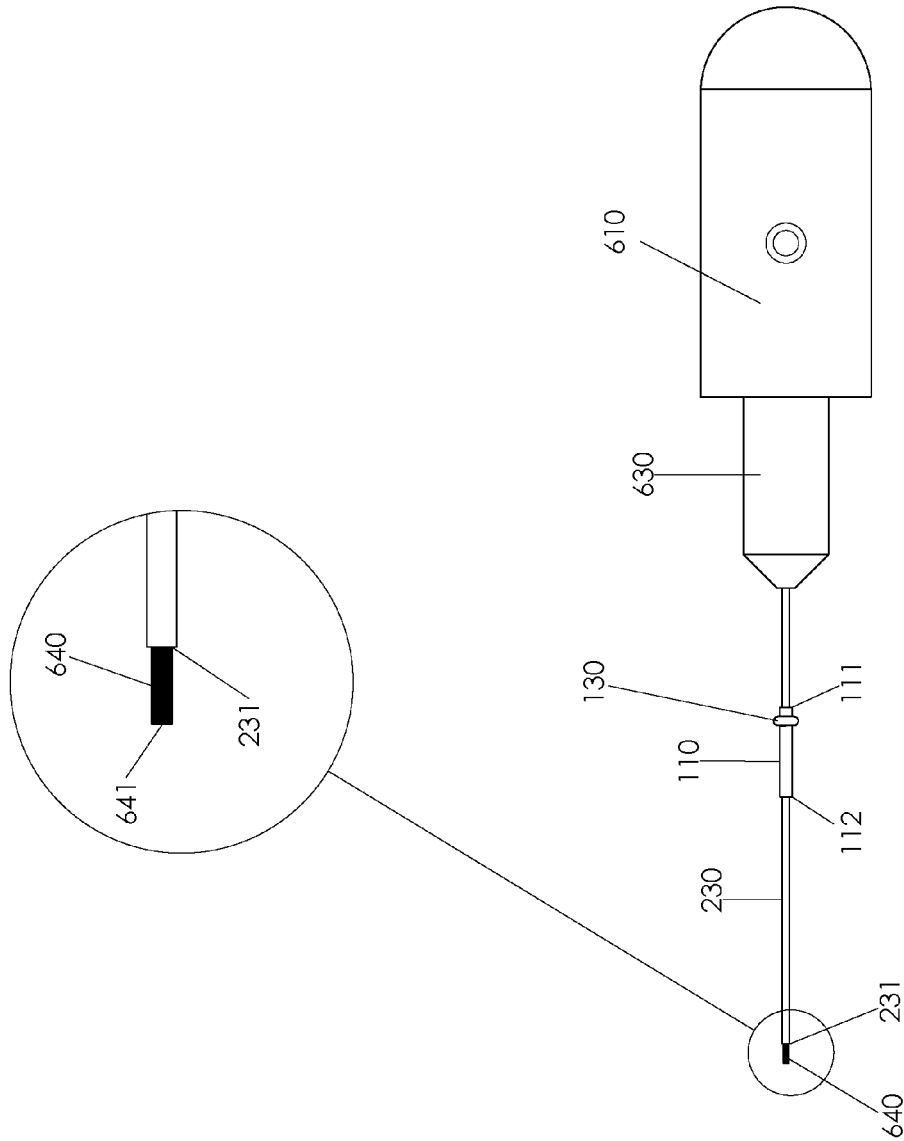


FIG. 8

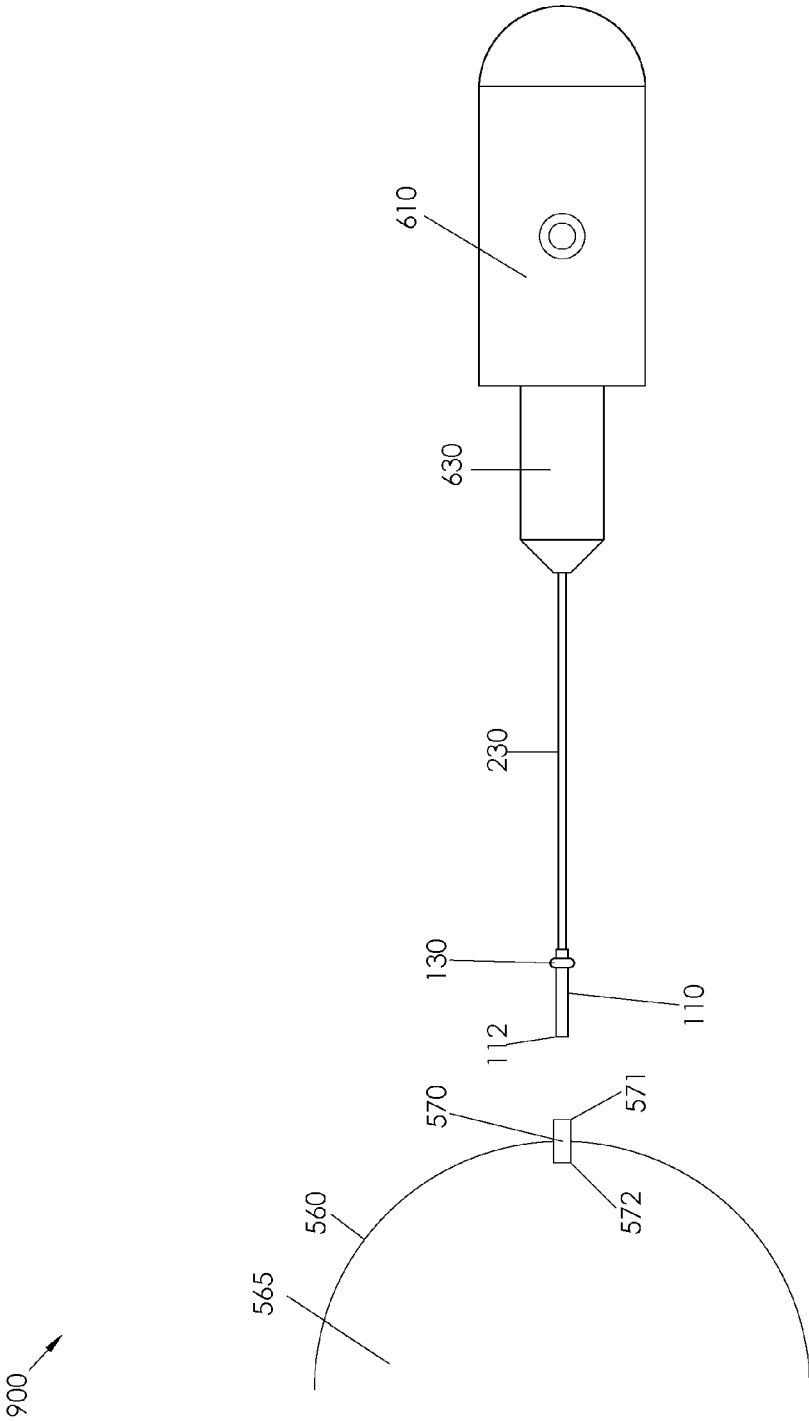


FIG. 9A

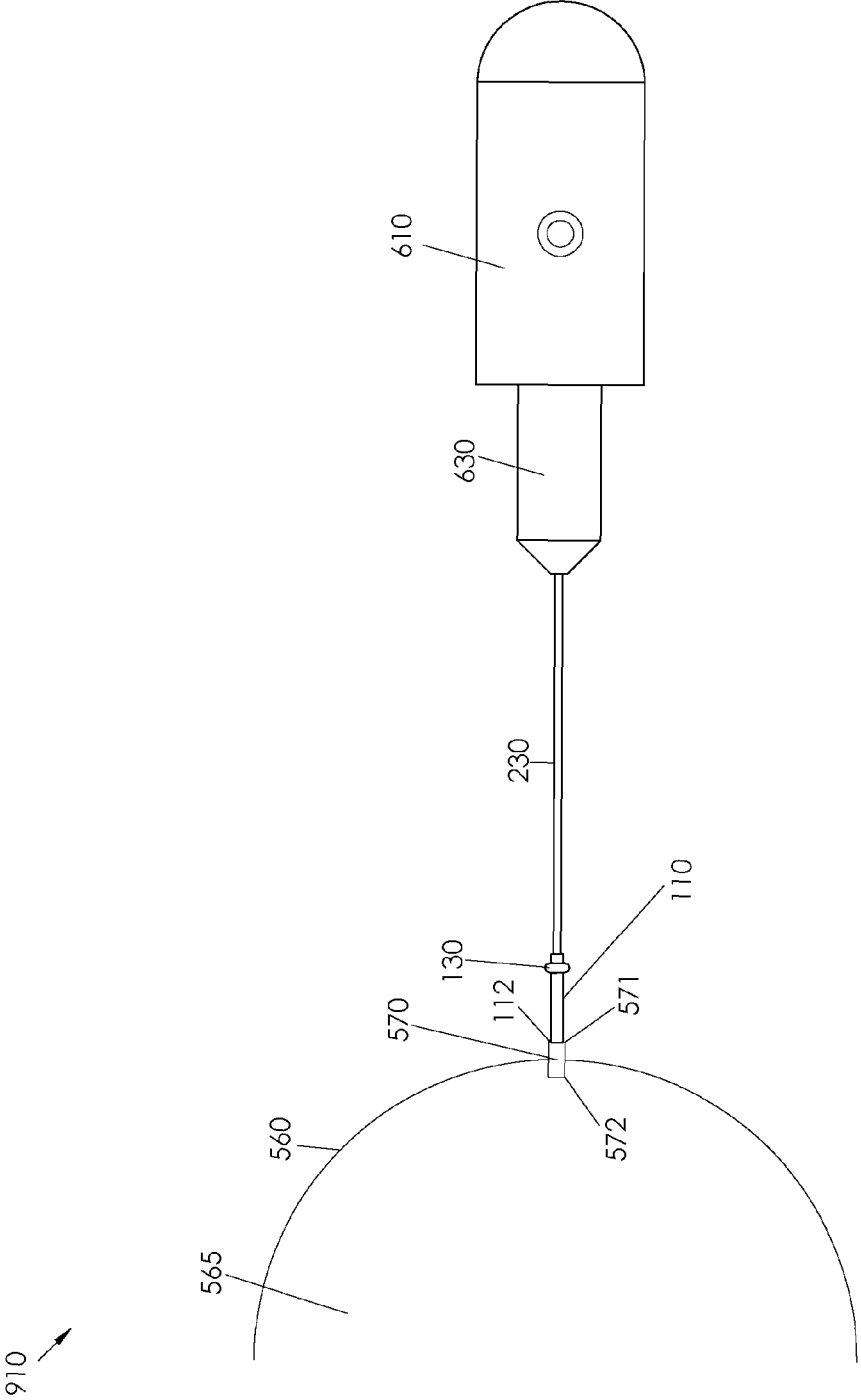


FIG. 9B

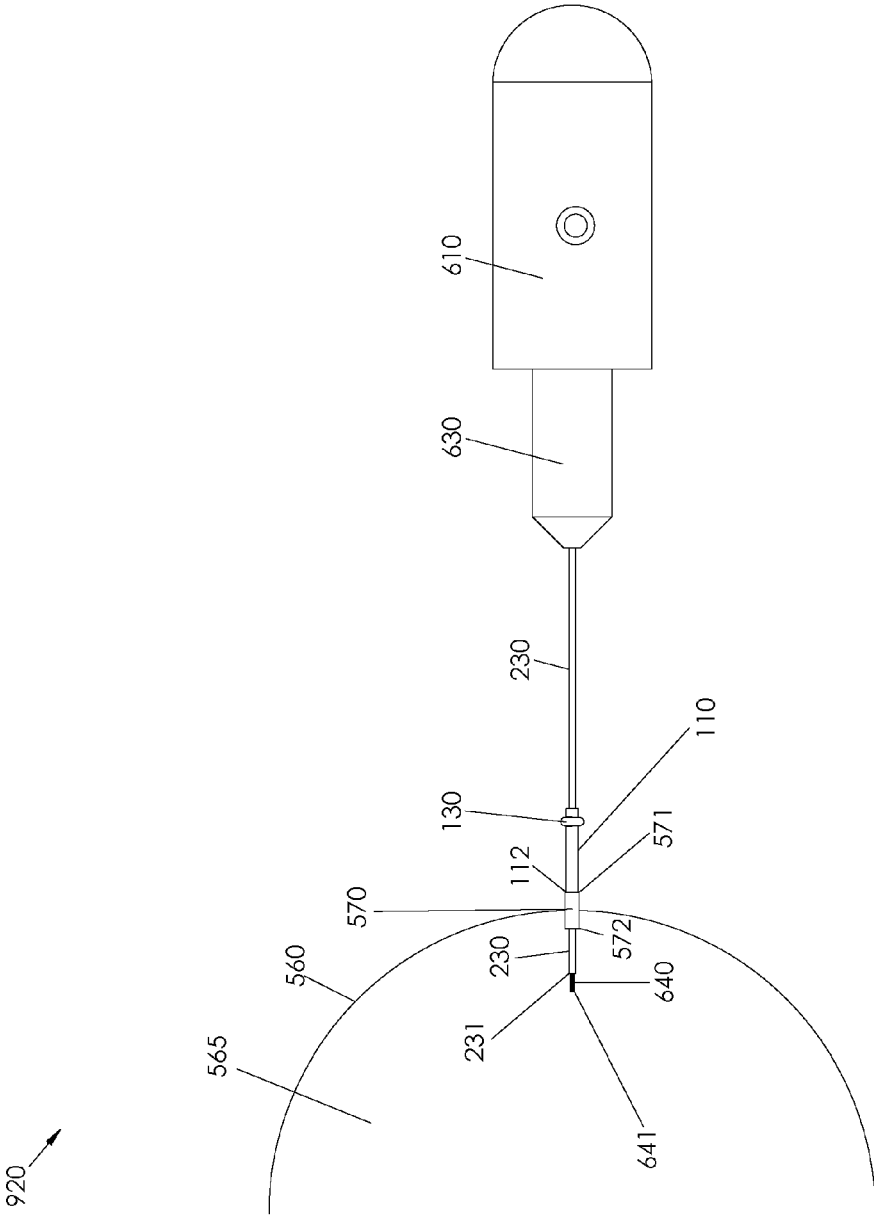


FIG. 9C

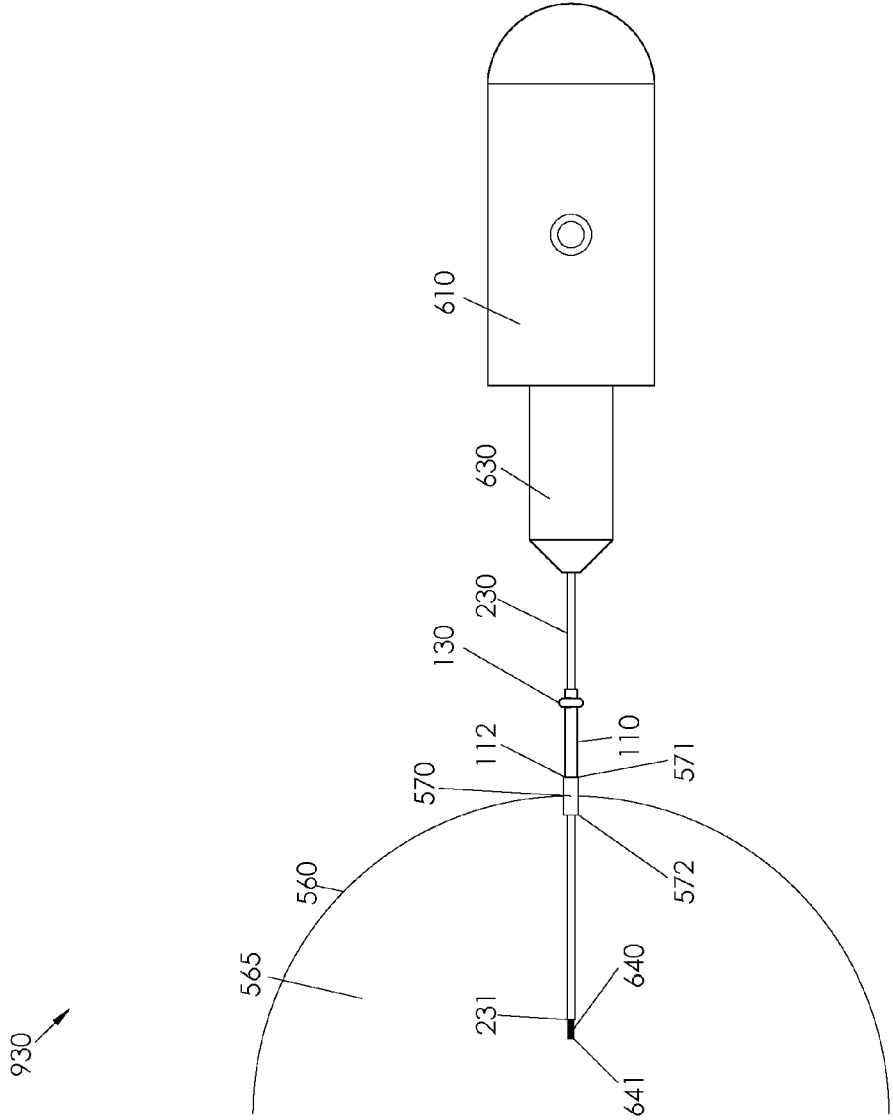


FIG. 9D



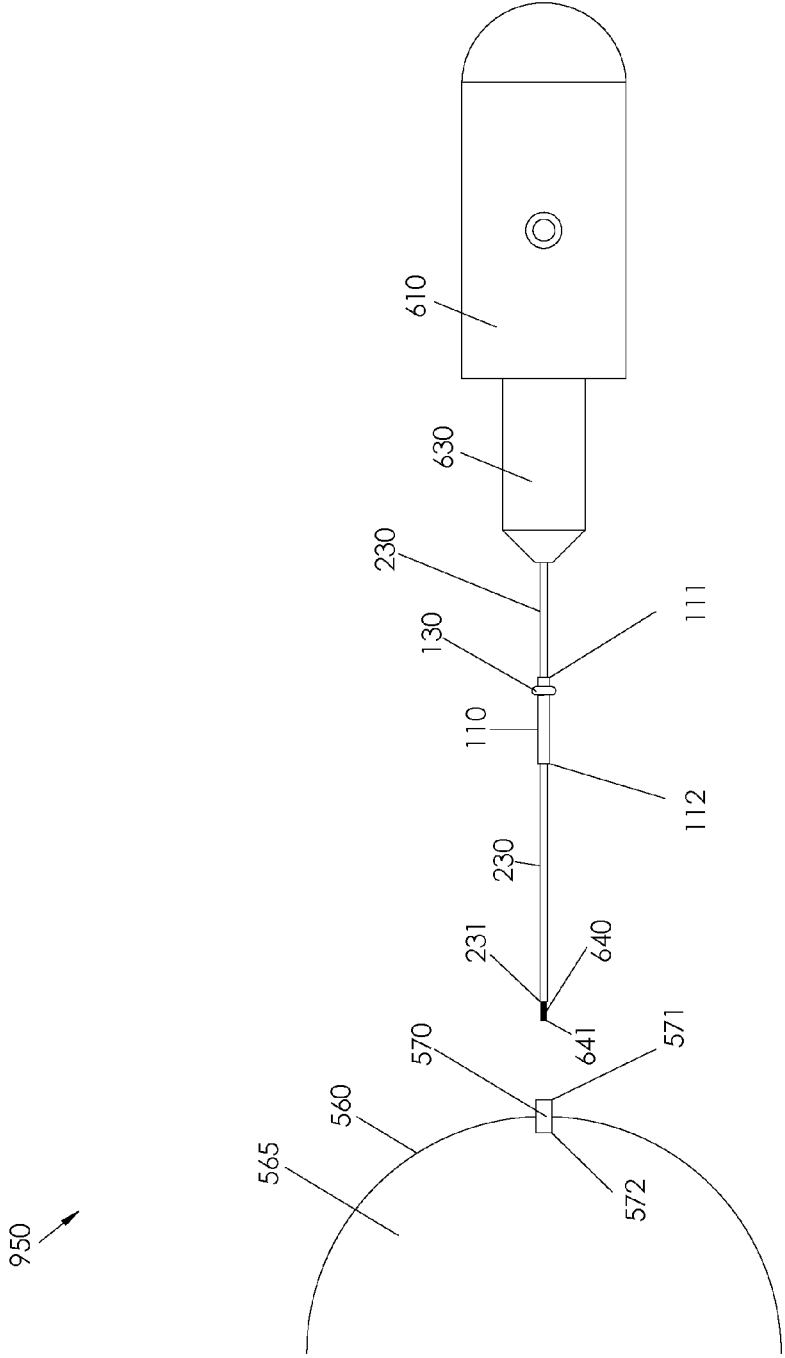


FIG. 9F



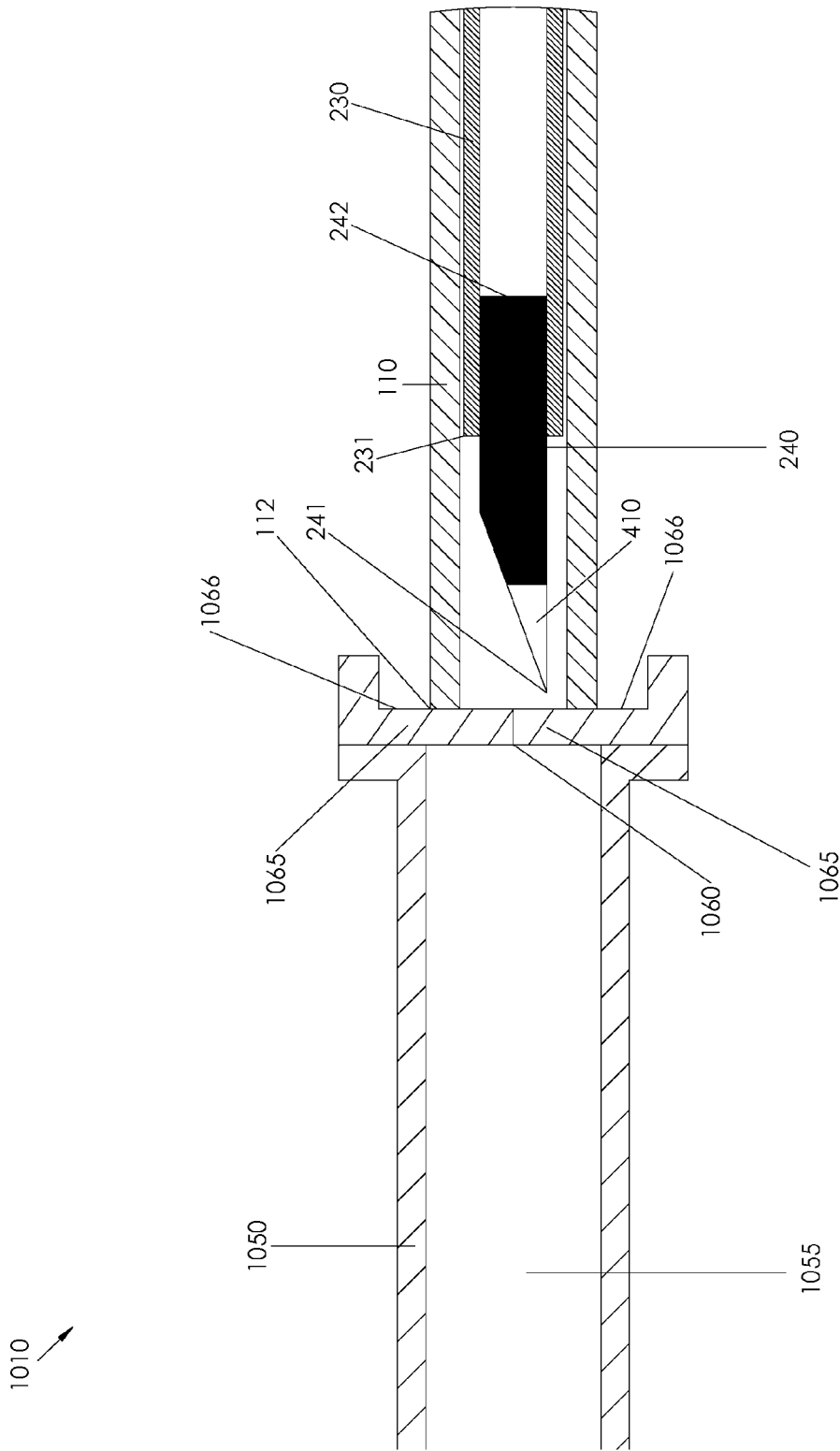


FIG. 10B

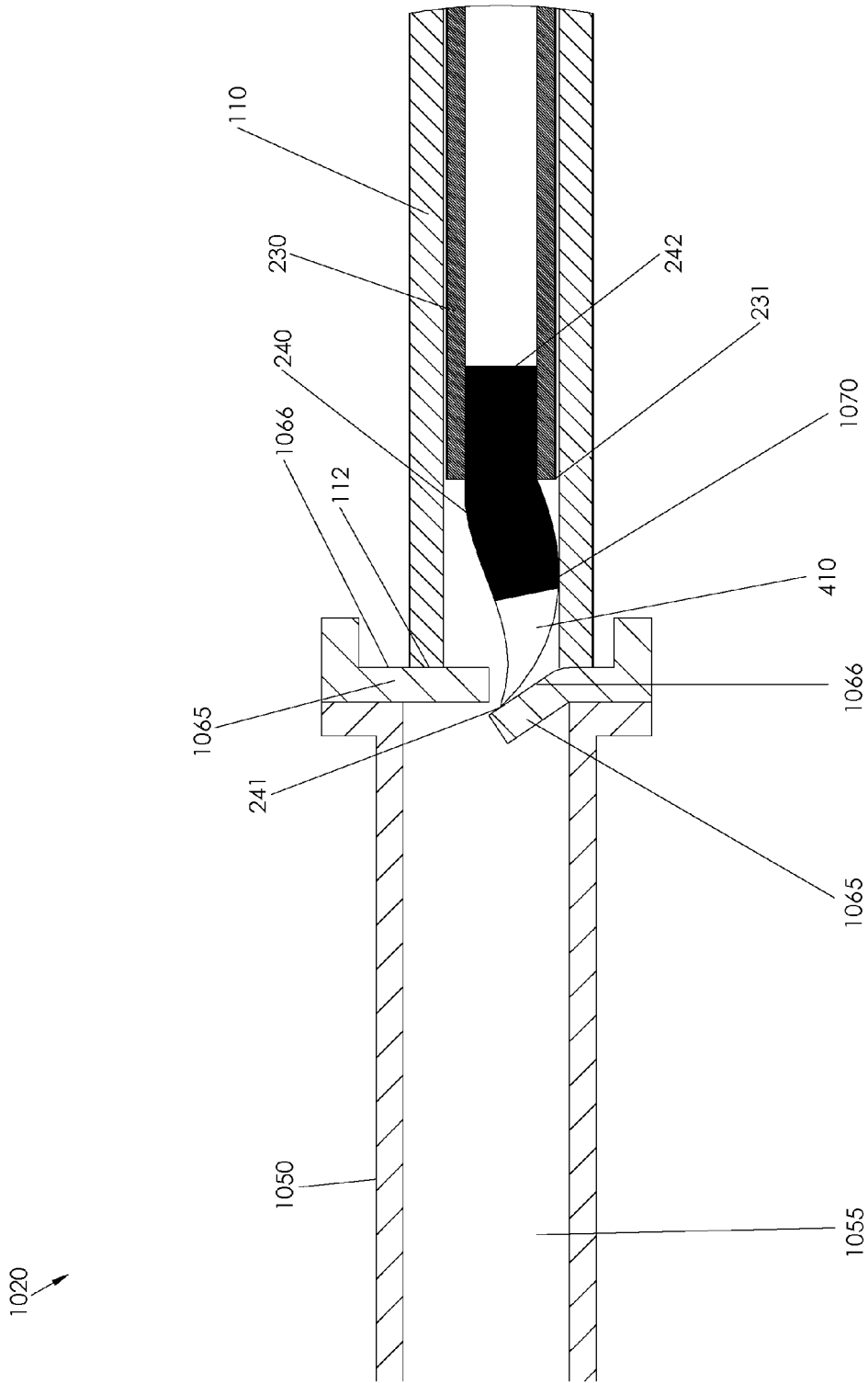


FIG. 10C

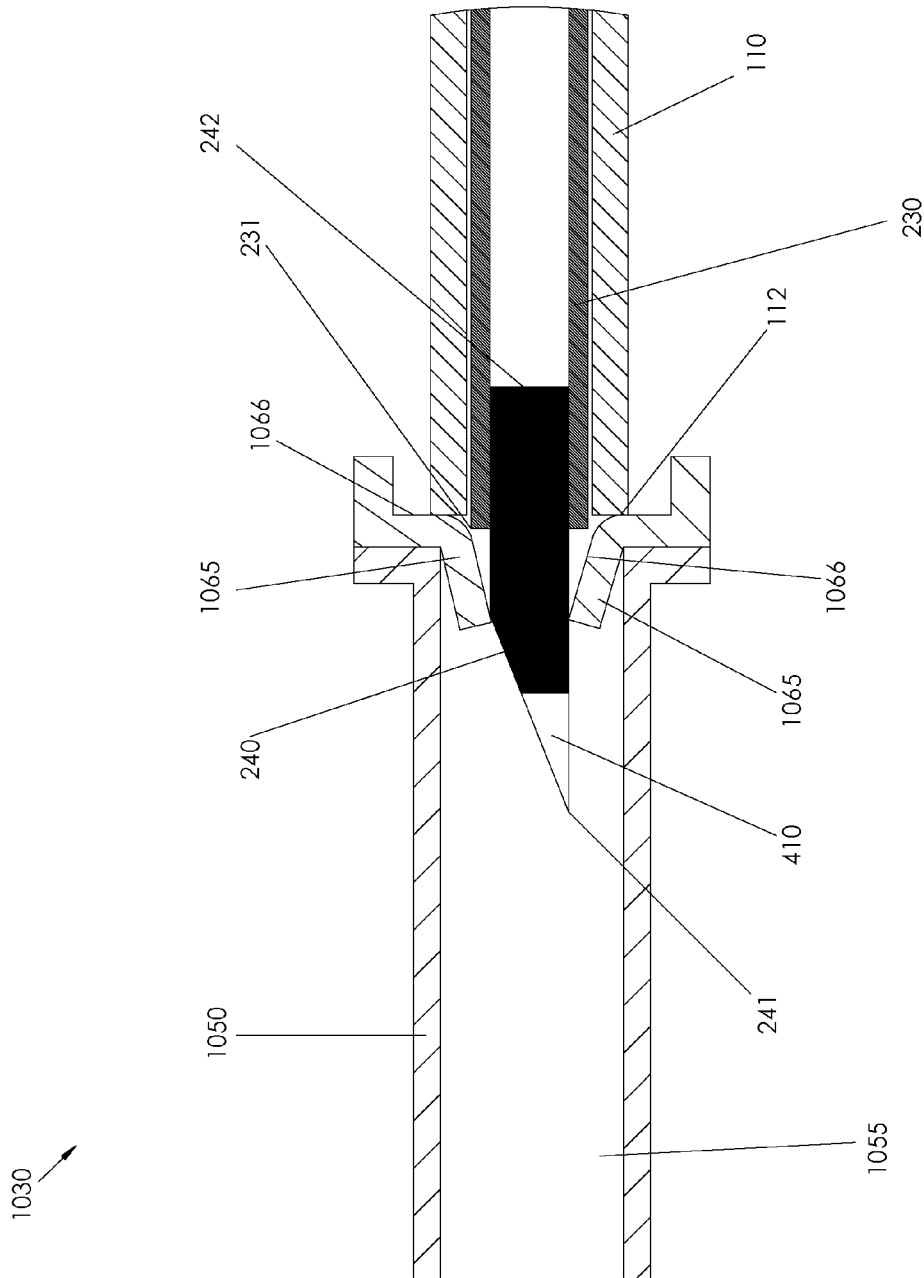


FIG. 10D

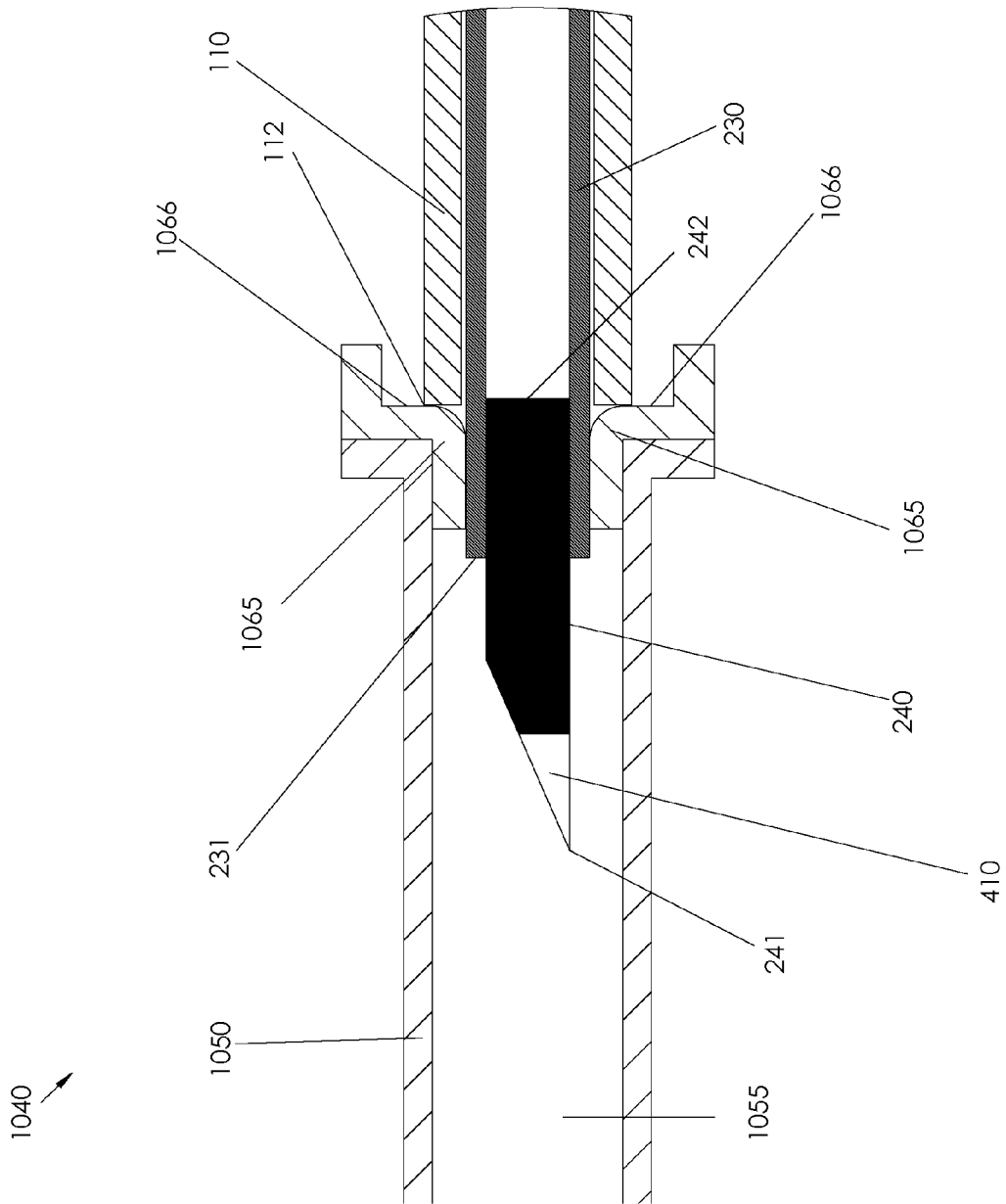


FIG. 10E



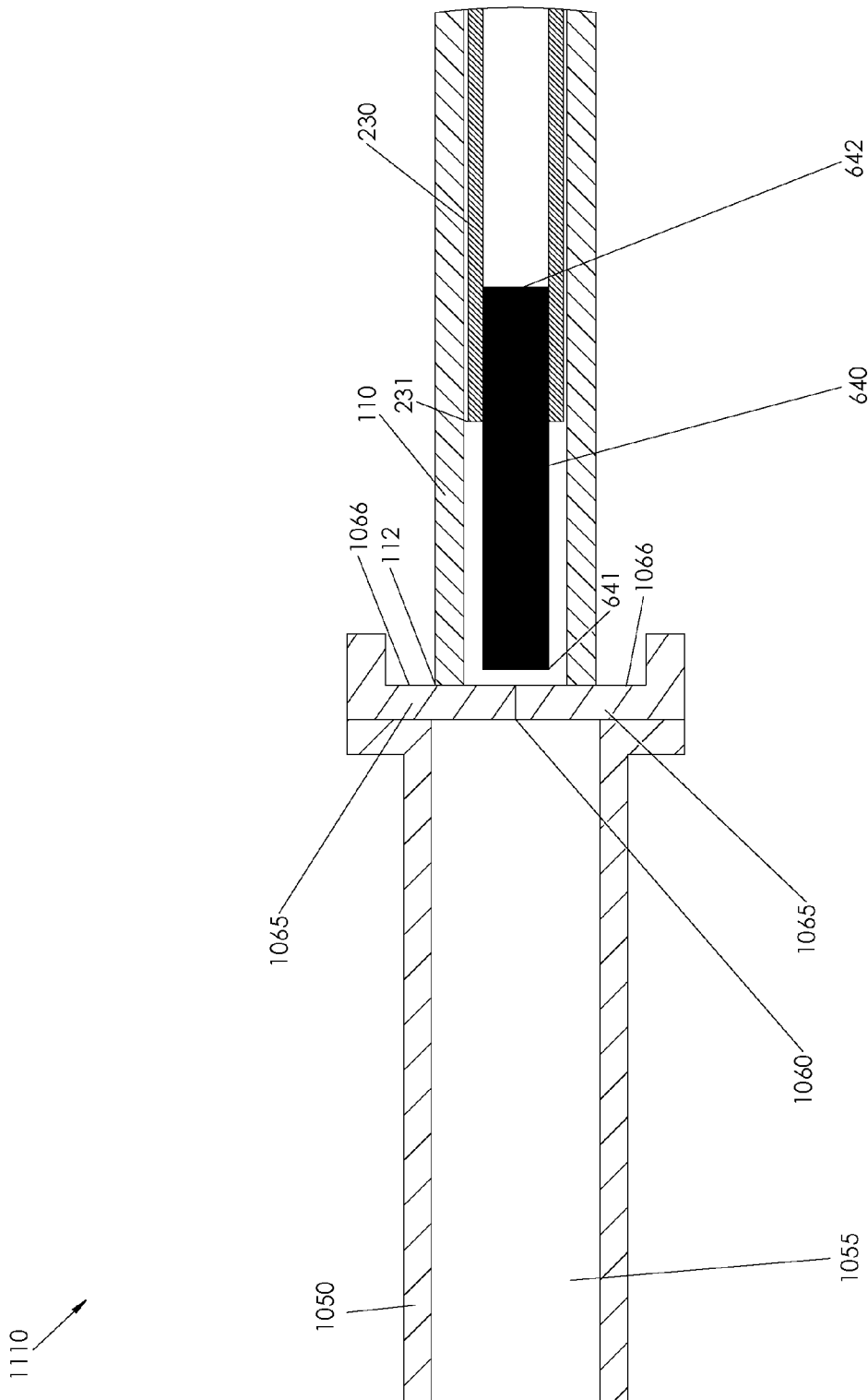


FIG. 11B

1120 ↗

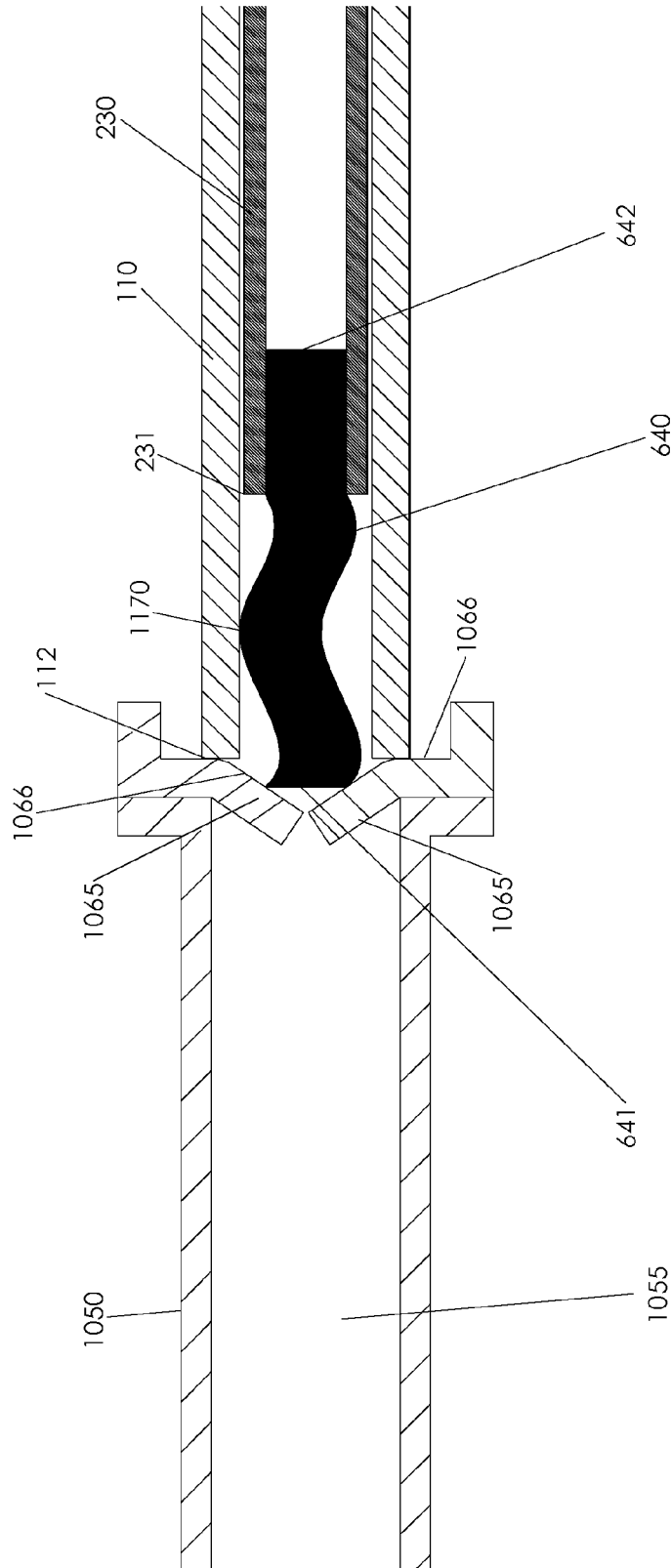


FIG. 11C

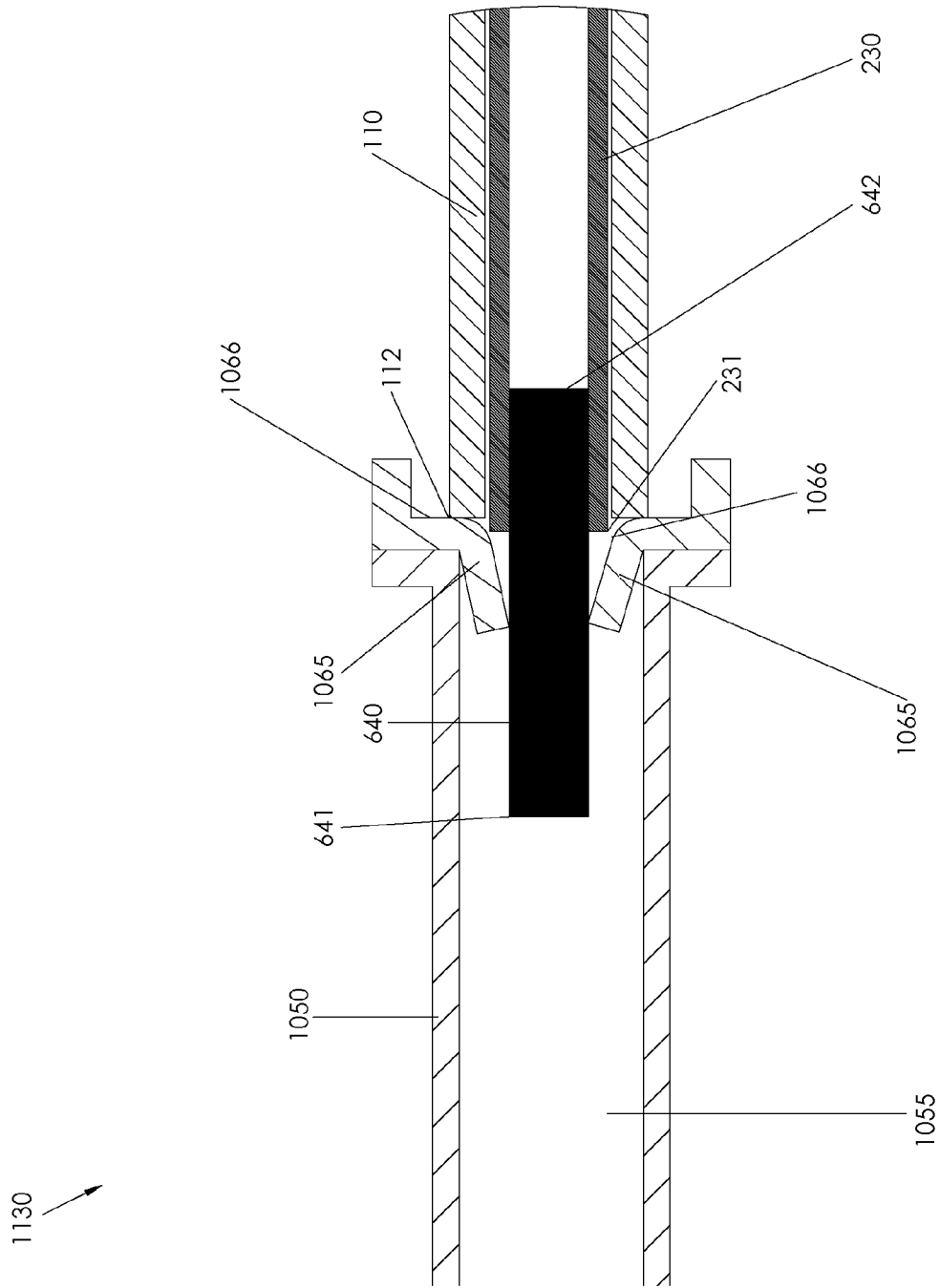


FIG. 11D

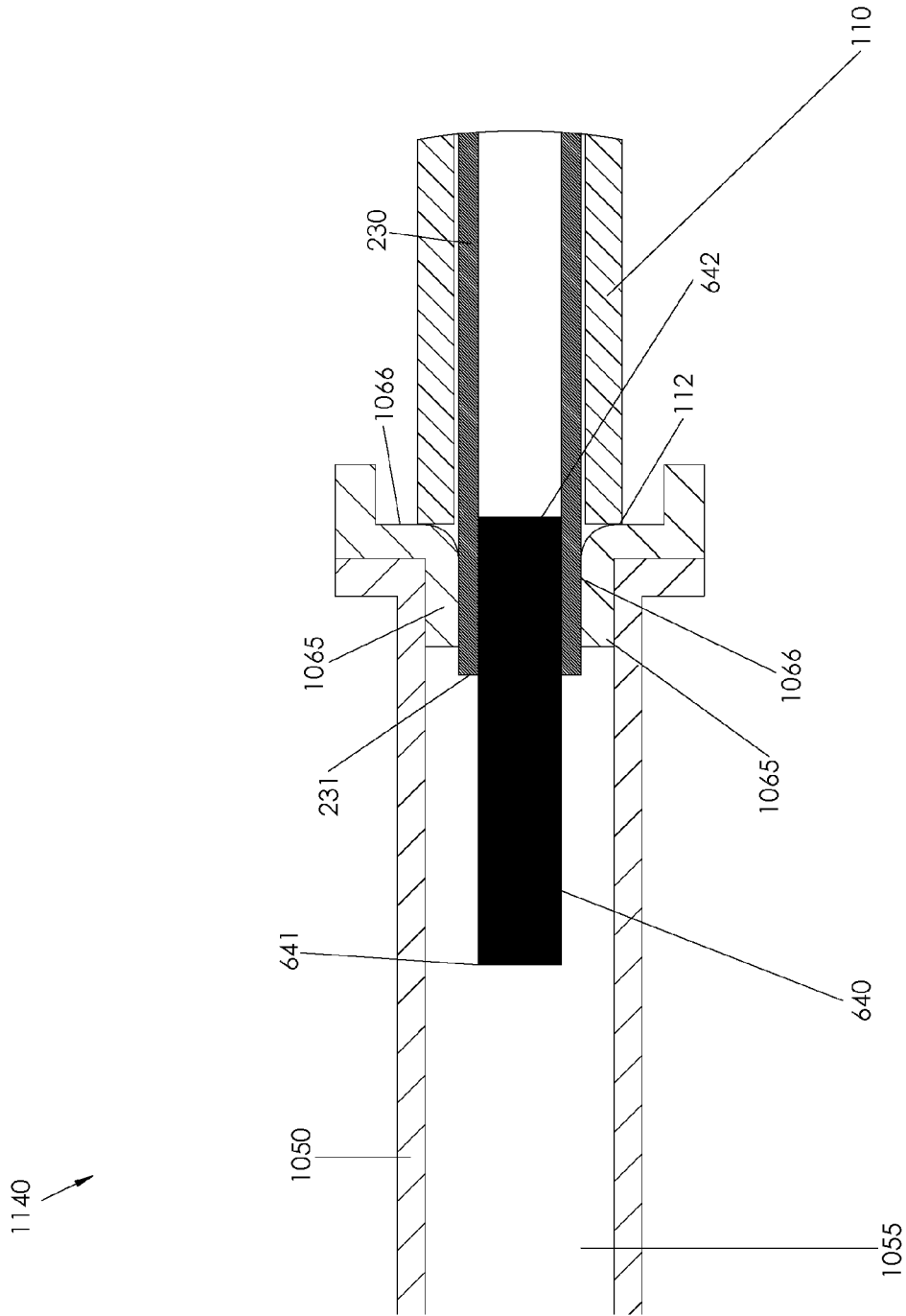


FIG. 11E

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**CANNULA INGRESS SYSTEM****CROSS-REFERENCE TO RELATED APPLICATIONS**

This Application claims the benefit of U.S. Provisional Application No. 62/062,678, Oct. 10, 2014.

**FIELD OF THE INVENTION**

The present disclosure relates to a medical device, and, more particularly, to a surgical instrument.

**BACKGROUND OF THE INVENTION**

Many ophthalmic surgical procedures in the posterior segment are performed through a cannula. For example, a surgeon may make an incision in the pars plana and then insert a cannula in the incision. The surgeon may then access the inner eye via the cannula. Ophthalmic surgical instruments having a soft, flexible tip may be difficult to insert into a cannula because the soft, flexible tip may deform as a surgeon attempts to ingress the cannula. For example, as a surgeon approaches a cannula to attempt a cannula ingress, a soft, flexible tip of an instrument must be precisely aligned with a cannula opening to prevent the soft, flexible tip from deforming.

**BRIEF SUMMARY OF THE INVENTION**

A cannula ingress system is presented. Illustratively, a cannula ingress system may comprise a tip stabilization mechanism having a tip stabilization mechanism distal end and a tip stabilization mechanism proximal end, a fixation mechanism, a hypodermic tube having a hypodermic tube distal end and a hypodermic tube proximal end, and a tip having a tip distal end and a tip proximal end. In one or more embodiments, the tip may be disposed within the hypodermic tube wherein the tip distal end extends from the hypodermic tube distal end. Illustratively, the fixation mechanism may be disposed within a fixation mechanism channel of the tip stabilization mechanism. In one or more embodiments, the tip stabilization mechanism may be disposed over the tip and the hypodermic tube distal end.

**BRIEF DESCRIPTION OF THE DRAWINGS**

The above and further advantages of the present invention may be better understood by referring to the following description in conjunction with the accompanying drawings in which like reference numerals indicate identical or functionally similar elements:

FIGS. 1A and 1B are schematic diagrams illustrating a tip stabilization system assembly;

FIG. 2 is a schematic diagram illustrating an exploded view of an instrument assembly;

FIGS. 3A and 3B are schematic diagrams illustrating an assembled instrument;

FIG. 4 is a schematic diagram illustrating an exposed membrane removing tip;

FIGS. 5A, 5B, 5C, 5D, 5E, and 5F are schematic diagrams illustrating a portion of a surgical procedure;

FIG. 6 is a schematic diagram illustrating an exploded view of an irrigating and aspirating instrument assembly;

FIGS. 7A and 7B are schematic diagrams illustrating an assembled irrigating and aspirating instrument;

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FIG. 8 is a schematic diagram illustrating an exposed irrigating and aspirating tip;

FIGS. 9A, 9B, 9C, 9D, 9E, and 9F are schematic diagrams illustrating a portion of a surgical procedure;

FIGS. 10A, 10B, 10C, 10D, and 10E are schematic diagrams illustrating a portion of a surgical procedure;

FIGS. 11A, 11B, 11C, 11D, and 11E are schematic diagrams illustrating a portion of a surgical procedure.

**DETAILED DESCRIPTION OF AN ILLUSTRATIVE EMBODIMENT**

FIGS. 1A and 1B are schematic diagrams illustrating a tip stabilization system assembly **100**. FIG. 1A illustrates a side view of a tip stabilization system assembly **100**. In one or more embodiments, tip stabilization system assembly **100** may comprise a tip stabilization mechanism **110** and a fixation mechanism **130**. Illustratively, tip stabilization mechanism **110** may comprise a tip stabilization mechanism distal end **111**, a tip stabilization mechanism proximal end **112**, a tip stabilization mechanism outer diameter **113**, a tip stabilization mechanism inner diameter **114**, and a fixation mechanism channel **115**. In one or more embodiments, fixation mechanism channel **115** may have a width in a range of 0.01 to 0.05 inches, e.g., fixation mechanism channel **115** may have a width of 0.033 inches. Illustratively, fixation mechanism channel **115** may have a width of less than 0.01 inches or greater than 0.05 inches. In one or more embodiments, tip stabilization mechanism outer diameter **113** may be a distance in a range of 0.01 to 0.04 inches, e.g., tip stabilization mechanism outer diameter **113** may be a distance of 0.0226 inches. Illustratively, tip stabilization mechanism outer diameter **113** may be a distance of less than 0.01 inches or greater than 0.04 inches. In one or more embodiments, tip stabilization mechanism inner diameter **114** may be a distance in a range of 0.005 to 0.039 inches, e.g., tip stabilization mechanism inner diameter **114** may be a distance of 0.02 inches. Illustratively, tip stabilization mechanism inner diameter **114** may be a distance of less than 0.005 inches or greater than 0.039 inches. In one or more embodiments, tip stabilization mechanism **110** may have an overall length in a range of 0.2 to 0.4 inches, e.g., tip stabilization mechanism **110** may have an overall length of 0.3 inches. Illustratively, tip stabilization mechanism **110** may have an overall length of less than 0.2 inches or greater than 0.4 inches. In one or more embodiments, tip stabilization mechanism **110** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, fixation mechanism **130** may comprise a fixation mechanism outer diameter **135** and a fixation mechanism inner diameter **136**. In one or more embodiments, fixation mechanism outer diameter **135** may be greater than tip stabilization mechanism outer diameter **113**, e.g., fixation mechanism outer diameter **135** may be a first distance and tip stabilization mechanism outer diameter **113** may be a second distance wherein the first distance is greater than the second distance. Illustratively, fixation mechanism outer diameter **135** may be less than tip stabilization mechanism outer diameter **113**, e.g., fixation mechanism outer diameter **135** may be a first distance and tip stabilization mechanism outer diameter **113** may be a second distance wherein the first distance is less than the second distance. In one or more embodiments, fixation mechanism outer diameter **135** may be equal to tip stabilization mechanism outer diameter **113**, e.g., fixation mechanism outer diameter **135** may be a first distance and tip stabilization mechanism outer diameter **113** may be a

second distance wherein the first distance is equal to the second distance. Illustratively, fixation mechanism outer diameter 135 may be greater than tip stabilization mechanism inner diameter 114, e.g., fixation mechanism outer diameter 135 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is greater than the second distance. In one or more embodiments, fixation mechanism outer diameter 135 may be less than tip stabilization mechanism inner diameter 114, e.g., fixation mechanism outer diameter 135 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is less than the second distance. Illustratively, fixation mechanism outer diameter 135 may be equal to tip stabilization mechanism inner diameter 114, e.g., fixation mechanism outer diameter 135 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is equal to the second distance. In one or more embodiments, fixation mechanism inner diameter 136 may be greater than tip stabilization mechanism outer diameter 113, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism outer diameter 136 may be a second distance where in the first distance is greater than the second distance. Illustratively, fixation mechanism inner diameter 136 may be less than tip stabilization mechanism outer diameter 113, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism outer diameter 136 may be a second distance where in the first distance is less than the second distance. In one or more embodiments, fixation mechanism inner diameter 136 may be equal to tip stabilization mechanism outer diameter 113, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism outer diameter 136 may be a second distance where in the first distance is equal to the second distance. Illustratively, fixation mechanism inner diameter 136 may be greater than tip stabilization mechanism inner diameter 114, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is greater than the second distance. In one or more embodiments, fixation mechanism inner diameter 136 may be less than tip stabilization mechanism inner diameter 114, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is less than the second distance. Illustratively, fixation mechanism inner diameter 136 may be equal to tip stabilization mechanism inner diameter 114, e.g., fixation mechanism inner diameter 136 may be a first distance and tip stabilization mechanism inner diameter 114 may be a second distance wherein the first distance is equal to the second distance. In one or more embodiments, fixation mechanism 130 may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. FIG. 1B illustrates a cross-sectional view of a tip stabilization system assembly 100. In one or more embodiments, fixation mechanism 130 may have a cross-sectional radius in a range of 0.005 to 0.03 inches, e.g., fixation mechanism 130 may have a cross-sectional radius of 0.015 inches. Illustratively, fixation mechanism 130 may have a cross-sectional radius of less than 0.005 inches or greater than 0.03 inches. In one or more embodiments, tip stabilization mechanism 110 may comprise an inner bore 120 having an inner bore distal end 121 and an inner bore proximal end 122. Illustratively, fixation mechanism 130 and tip stabilization mechanism 110 may be

manufactured as a single component, e.g., fixation mechanism 130 may comprise a feature of tip stabilization mechanism 110. In one or more embodiments, tip stabilization mechanism 110 may comprise a tube and fixation mechanism 130 may comprise a discontinuity in inner bore 120, e.g., tube fixation mechanism 130 may comprise a portion of inner bore 120 having an inner diameter less than tip stabilization mechanism inner diameter 114.

FIG. 2 is a schematic diagram illustrating an exploded view of an instrument assembly 200. In one or more embodiments, an instrument assembly 200 may comprise a tip stabilization mechanism 110, a fixation mechanism 130, a handle 210, an identification ring 220, a hypodermic tube 230, and a membrane removing tip 240. Illustratively, handle 210 may comprise a handle distal end 211, a handle proximal end 212, an identification ring channel 215, a handle grip 216, and a hypodermic tube interface 217. In one or more embodiments, handle 210 may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, hypodermic tube 230 may comprise a hypodermic tube distal end 231 and a hypodermic tube proximal end 232. In one or more embodiments, hypodermic tube 230 may be manufactured at dimensions configured for performing microsurgical procedures, e.g., hypodermic tube 230 may be manufactured at dimensions configured for performing ophthalmic surgical procedures. Illustratively, hypodermic tube 230 may be 20 gauge, 23 gauge, 25 gauge, 27 gauge, etc. In one or more embodiments, hypodermic tube 230 may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, membrane removing tip 240 may comprise a membrane removing tip distal end 241 and a membrane removing tip proximal end 242. In one or more embodiments, membrane removing tip 240 may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, membrane removing tip 240 may be manufactured from silicone, e.g., biocompatible silicone.

FIGS. 3A and 3B are schematic diagrams illustrating an assembled instrument 300. FIG. 3A illustrates a side view of an assembled instrument 300. In one or more embodiments, fixation mechanism 130 may be disposed over a portion of tip stabilization mechanism 110, e.g., fixation mechanism 130 may be disposed between tip stabilization mechanism distal end 111 and tip stabilization mechanism proximal end 112. Illustratively, a portion of fixation mechanism 130 may be disposed within a portion of tip stabilization mechanism 110, e.g., a portion of fixation mechanism 130 may be disposed within fixation mechanism channel 115. In one or more embodiments, a portion of fixation mechanism 130 may be disposed within inner bore 120. Illustratively, a portion of fixation mechanism 130 may be fixed to a portion of tip stabilization mechanism 110, e.g., fixation mechanism inner diameter 136 may be fixed to a portion of tip stabilization mechanism 110. In one or more embodiments, a portion of fixation mechanism 130 may be fixed to a portion of tip stabilization mechanism 110 by a force of friction or by any suitable fixation means, e.g., a portion of fixation mechanism 130 may be fixed to a portion of tip stabilization mechanism 110 by an adhesive, a weld, etc.

Illustratively, tip stabilization mechanism 110 may be disposed over a portion of hypodermic tube 230, e.g., tip stabilization mechanism distal end 111 may be disposed over hypodermic tube distal end 231. In one or more embodiments, a portion of hypodermic tube 230 may be disposed

within inner bore **120**, e.g., hypodermic tube distal end **231** may be disposed within inner bore **120**. Illustratively, hypodermic tube **230** may be disposed within fixation mechanism **130**, e.g., hypodermic tube **230** may be disposed within fixation mechanism inner diameter **136**. In one or more embodiments, hypodermic tube **230** may not be disposed within fixation mechanism **130**, e.g., a portion of hypodermic tube **230** may be disposed adjacent to a portion of fixation mechanism **130**. Illustratively, fixation mechanism **130** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**, e.g., a static friction force between fixation mechanism **130** and hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**. In one or more embodiments, a contact between a portion of fixation mechanism **130** and a portion of hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**. Illustratively, fixation mechanism **130** may be fixed to both tip stabilization mechanism **110** and hypodermic tube **230** wherein a static friction force between fixation mechanism **130** and hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**.

Illustratively, a portion of membrane removing tip **240** may be disposed within hypodermic tube **230**, e.g., membrane removing tip proximal end **242** may be disposed within hypodermic tube distal end **231**. In one or more embodiments, a portion of membrane removing tip **240** may be disposed within hypodermic tube **230** wherein a portion of membrane removing tip **240** extends from a portion of hypodermic tube **230**, e.g., a portion of membrane removing tip **240** may be disposed within hypodermic tube **230** wherein membrane removing tip distal end **241** extends from hypodermic tube distal end **231**. Illustratively, a portion of membrane removing tip **240** may be fixed within hypodermic tube **230**, e.g., membrane removing tip proximal end **242** may be fixed within hypodermic tube **230**. In one or more embodiments, a portion of membrane removing tip **240** may be fixed within hypodermic tube **230** by an adhesive or any suitable fixation means, e.g., a portion of membrane removing tip **240** may be fixed within hypodermic tube **230** by a friction fit, a weld, etc.

Illustratively, identification ring **220** may be disposed over a portion of handle **210**, e.g., identification ring **220** may be disposed within identification ring channel **215**. In one or more embodiments, identification ring **220** may be fixed to a portion of handle **210**, e.g., identification ring **220** may be fixed within identification ring channel **215** by an adhesive, a force of friction, a weld, or any suitable fixation means. Illustratively, identification ring **220** may be configured to indicate one or more properties of assembled instrument **300** to a surgeon, a nurse, or a surgical technician, e.g., identification ring **220** may comprise a color or a marking configured to visually indicate a minimum cannula gauge size that hypodermic tube **230** may ingress to perform a surgical procedure. In one or more embodiments, identification ring **220** may be manufactured from a shape memory material, e.g., Nitinol. Illustratively, identification ring **220** may be manufactured from an elastomer material, e.g., a biocompatible elastomer material. In one or more embodiments, identification ring **220** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials.

FIG. 3B illustrates a cross-sectional view of an assembled instrument **300**. In one or more embodiments, an assembled

instrument **300** may comprise an inner chamber proximal taper **310**, an inner chamber **320**, an inner chamber distal taper **330**, and a hypodermic tube housing **340**. Illustratively, a portion of hypodermic tube **230** may be disposed within a portion of handle **210**, e.g., a portion of hypodermic tube **230** may be disposed within a portion of handle **210** wherein the portion of hypodermic tube **230** may be adjacent to hypodermic tube interface **117**. In one or more embodiments, a portion of hypodermic tube **230** may be disposed within hypodermic tube housing **340**, e.g., hypodermic tube proximal end **232** may be disposed within hypodermic tube housing **340**. In one or more embodiments, a portion of hypodermic tube **230** may be fixed within a portion of handle **210**, e.g., a portion of hypodermic tube **230** may be fixed within hypodermic tube housing **340**. Illustratively, a portion of hypodermic tube **230** may be fixed within hypodermic tube housing by an interference fit, an adhesive, a weld, etc. In one or more embodiments, a portion of hypodermic tube **230** may be fixed within a portion of handle **210** wherein hypodermic tube distal end **231** extends out from hypodermic tube housing **340**.

Illustratively, membrane removing tip **240** may be disposed within hypodermic tube **230** and tip stabilization mechanism **110**, e.g., membrane removing tip distal end **241** may be disposed between tip stabilization mechanism distal end **111** and tip stabilization mechanism proximal end **112**. In one or more embodiments, membrane removing tip **240** may be disposed within hypodermic tube **230** and tip stabilization mechanism **110** wherein membrane removing tip proximal end **242** may be disposed between hypodermic tube distal end **231** and hypodermic tube proximal end **232**. Illustratively, hypodermic tube **230** may be disposed within tip stabilization mechanism **110** wherein hypodermic tube distal end **231** is disposed between tip stabilization mechanism distal end **111** and tip stabilization mechanism proximal end **112**.

FIG. 4 is a schematic diagram illustrating an exposed membrane removing tip **400**. In one or more embodiments, membrane removing tip **240** may be colored black, e.g., to enhance visualization of membrane removing tip **240**. For example, membrane removing tip **240** may be manufactured from black silicone to enhance visualization of membrane removing tip **240**. Illustratively, membrane removing tip **240** may be colored any color not contained within Johannes Itten's color wheel to enhance visualization of membrane removing tip **240**, e.g., membrane removing tip **240** may be colored black or grey to enhance visualization of membrane removing tip **240**. In one or more embodiments, an exposed membrane removing tip **400** may comprise an abrasive surface **410** and a membrane removing tip taper angle **420**. Illustratively, membrane removing tip taper angle **420** may be an angle in a range of 10.0 to 60.0 degrees, e.g., membrane removing taper angle **420** may be an angle of 25.0 degrees. In one or more embodiments, membrane removing taper angle **420** may be an angle of less than 10.0 degrees or greater than 60.0 degrees.

In one or more embodiments, abrasive surface **410** may be configured to grasp a portion of a membrane, e.g., abrasive surface **410** may be configured to grasp a portion of an internal limiting membrane. Illustratively, a surgeon may maneuver a portion of abrasive surface **410** across a portion of a membrane, e.g., to raise a portion of the membrane. In one or more embodiments, abrasive surface **410** may be configured to grasp a portion of a first tissue disposed over a second tissue without damaging the second tissue. Illustratively, abrasive surface **410** may be configured to grasp a first tissue having a convex surface geometry disposed over

a second tissue having a convex surface geometry without damaging the second tissue. In one or more embodiments, abrasive surface **410** may be manufactured by fixing particles, e.g., inert particles, to a portion of membrane removing tip **240**. Illustratively, particles may be fixed to a portion of membrane removing tip **240**, e.g., by an adhesive or any suitable fixation means. In one or more embodiments, particles may be fixed to a portion of membrane removing tip **240** by a biocompatible high temperature epoxy. Illustratively, particles may be fixed to a portion of membrane removing tip **240** by a biocompatible spectrally transparent epoxy. In one or more embodiments, a portion of membrane removing tip **240** may be coated by a material configured to facilitate adhesion of particles. Illustratively, a portion of membrane removing tip **240** may be coated by a material, e.g., silicon, and then particles may be fixed to the material, e.g., by an adhesive or any suitable fixation means. In one or more embodiments, abrasive surface **410** may be manufactured by fixing particles to a portion of membrane removing tip **240**, e.g., particles may comprise diamond particles, sapphire particles, ruby particles, emerald particles, etc. Illustratively, abrasive surface **410** may be manufactured by fixing biocompatible particles to a portion of membrane removing tip **240**. In one or more embodiments, abrasive surface **410** may be manufactured by fixing particles having particle diameters in a range of 5.0 to 25.0 micrometers to a portion of membrane removing tip **240**, e.g., abrasive surface **410** may be manufactured by fixing particles having particle diameters of 15.0 micrometers to a portion of membrane removing tip **240**. Illustratively, abrasive surface **410** may be manufactured by fixing particles having particle diameters less than 5.0 micrometers or greater than 25.0 micrometers to a portion of membrane removing tip **240**. In one or more embodiments, membrane removing tip **240** may be colored black, e.g., to enhance visualization of abrasive surface **410**. For example, membrane removing tip **240** may be manufactured from black silicone to enhance visualization of abrasive surface **410**. Illustratively, membrane removing tip **240** may be colored any color not contained within Johannes Itten's color wheel to enhance visualization of abrasive surface **410**, e.g., membrane removing tip **240** may be colored black or grey to enhance visualization of abrasive surface **410**.

In one or more embodiments, abrasive surface **410** may be manufactured by modifying membrane removing tip **240**, e.g., by an electric discharge machine. Illustratively, abrasive surface **410** may be manufactured by actuating a portion of membrane removing tip **240** relative to a wire of an electric discharge machine, e.g., to form a plurality of micropillars. In one or more embodiments, abrasive surface **410** may be manufactured by actuating a wire of an electric discharge machine relative to a portion of membrane removing tip **240**, e.g., to form a plurality of micropillars. Illustratively, membrane removing tip **240** may be modified, e.g., by an electric discharge machine, wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars. Illustratively, abrasive surface **410** may be manufactured by modifying membrane removing tip **240**, e.g., by laser ablation. In one or more embodiments, abrasive surface **410** may be manufactured by modifying membrane removing tip **240**, e.g., by femtosecond laser ablation. Illustratively, abrasive surface **410** may be manufactured by applying laser energy to a portion of membrane removing tip **240** wherein the laser energy is applied in geometric patterns configured to fabricate micropillars on a surface of membrane removing tip **240**, e.g., the laser energy may be applied in concentric circles, polygons, etc. In one or more embodi-

ments, abrasive surface **410** may be manufactured by applying laser energy to a portion of membrane removing tip **240** wherein the laser energy is applied repeatedly in geometric patterns configured to fabricate micropillars on a surface of membrane removing tip **240**, e.g., the laser energy may be repeatedly applied in concentric circles, polygons, etc. Illustratively, membrane removing tip **240** may be modified, e.g., by laser ablation, wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars. In one or more embodiments, membrane removing tip **240** may be modified wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars and then membrane removing tip **240** may be modified to manufacture membrane removing tip **240**. Illustratively, one or more portions of membrane removing tip **240** may comprise a plurality of micropillars. In one or more embodiments, membrane removing tip **240** may be modified, e.g., by an electric discharge machine, to manufacture membrane removing tip **240** and then membrane removing tip **240** may be modified, e.g., by laser ablation, wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars.

Illustratively, abrasive surface **410** may be manufactured by modifying membrane removing tip **240**, e.g., by deep reactive-ion etching. In one or more embodiments, abrasive surface **410** may be manufactured by modifying membrane removing tip **240**, e.g., by the Bosch process of time-multiplexed etching. Illustratively, abrasive surface **410** may be manufactured by exposing a portion of membrane removing tip **240** to repeated cycles of isotropic plasma etching followed by deposition of a chemically inert passivation layer to fabricate a plurality of micropillars on a surface of membrane removing tip **240**. In one or more embodiments, abrasive surface **410** may be manufactured by fabricating a plurality of micropillars on a substrate and then fixing the substrate to a portion of membrane removing tip **240**. Illustratively, membrane removing tip **240** may be modified, e.g., by deep reactive-ion etching, wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars. In one or more embodiments, membrane removing tip **240** may be modified wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars and then membrane removing tip **240** may be modified to manufacture membrane removing tip **240**. Illustratively, one or more portions of membrane removing tip **240** may comprise a plurality of micropillars. In one or more embodiments, membrane removing tip **240** may be modified, e.g., by an electric discharge machine, to manufacture membrane removing tip **240** and then membrane removing tip **240** may be modified, e.g., by deep reactive-ion etching, wherein one or more portions of membrane removing tip **240** comprise a plurality of micropillars.

Illustratively, abrasive surface **410** may comprise a plurality of micropillars, e.g., abrasive surface **410** may comprise one or more micropillar arrays. In one or more embodiments, abrasive surface **410** may comprise a plurality of micropillars having micropillar diameters in a range of 5.0 to 25.0 micrometers, e.g., abrasive surface **410** may comprise a plurality of micropillars having micropillar diameters of 15.0 micrometers. In one or more embodiments, abrasive surface **410** may comprise a plurality of micropillars having micropillar diameters less than 5.0 micrometers or greater than 25.0 micrometers. Illustratively, abrasive surface **410** may comprise a plurality of micropillars having micropillar heights in a range of 0.25 to 3.0 micrometers, e.g., abrasive surface **410** may comprise a plurality of micropillars having micropillar heights of 2.25 micrometers. In one or more

embodiments, abrasive surface 410 may comprise a plurality of micropillars having micropillar heights less than 0.25 micrometers or greater than 3.0 micrometers. Illustratively, abrasive surface 410 may comprise a plurality of micropillars having micropillar heights in a range of 10.0 to 95.0 percent of the average thickness of an internal limiting membrane, e.g., abrasive surface 410 may comprise a plurality of micropillars having micropillar heights of 80.0 percent of the average thickness of an internal limiting membrane. In one or more embodiments, abrasive surface 410 may comprise a plurality of micropillars having micropillar orientations normal to a portion of a surface of membrane removing tip 240. Illustratively, abrasive surface 410 may comprise a plurality of micropillars having micropillar orientations at an angle relative to a portion of a surface of membrane removing tip 240. In one or more embodiments, abrasive surface 410 may comprise a plurality of micropillars having micropillar orientations at an angle in a range of 60.0 to 89.0 degrees relative to a portion of a surface of membrane removing tip 240, e.g., abrasive surface 410 may comprise a plurality of micropillars having micropillar orientations at an angle of 85.0 degrees relative to a portion of a surface of membrane removing tip 240. Illustratively, abrasive surface 410 may comprise a plurality of micropillars having micropillar orientations at an angle less than 60.0 degrees or greater than 89.0 degrees relative to a portion of a surface of membrane removing tip 240.

FIGS. 5A, 5B, 5C, 5D, 5E, and 5F are schematic diagrams illustrating a portion of a surgical procedure. FIG. 5A illustrates a cannula approach 500. Illustratively, a surgeon may maneuver an assembled instrument 300 towards an eye 560 to perform a portion of a surgical procedure, e.g., a surgeon may maneuver an assembled instrument 300 towards a cannula 570 to perform a portion of a surgical procedure. In one or more embodiments, cannula 570 may comprise a cannula distal end 571 and a cannula proximal end 572. Illustratively, cannula 570 may be disposed within an incision in eye 560, e.g., cannula 570 may be disposed within an incision in eye 560 wherein cannula distal end 571 extends out from an outer surface of eye 560 and cannula proximal end 572 is disposed within an inner eye 565. In one or more embodiments, a cannula approach 500 may comprise an attempt to guide tip stabilization mechanism proximal end 112 towards cannula distal end 571.

FIG. 5B illustrates a cannula contact 510. Illustratively, a surgeon may approach cannula 570 with assembled instrument 300 until a portion of assembled instrument 300 contacts a portion of cannula 570, e.g., a surgeon may approach cannula 570 with assembled instrument 300 until tip stabilization mechanism proximal end 112 contacts cannula distal end 571. In one or more embodiments, a cannula contact 510 may comprise a contact between tip stabilization mechanism proximal end 112 and cannula distal end 571. Illustratively, a cannula contact 510 may be configured to apply a force to fixation mechanism 130, e.g., a cannula contact 510 may apply a normal force to tip stabilization mechanism 110. In one or more embodiments, an application of a normal force to tip stabilization mechanism 110 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, an application of a normal force to tip stabilization mechanism 110 may not be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip

stabilization mechanism 110 may be less than a static friction force between fixation mechanism 130 and hypodermic tube 230.

FIG. 5C illustrates a partial membrane removing tip insertion 520. Illustratively, a surgeon may guide a portion of assembled instrument 300 into cannula 570, e.g., by advancing handle 210 towards eye 560 after a cannula contact 510. In one or more embodiments, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism distal end 111 towards handle distal end 211. In one or more embodiments, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to apply a force to tip stabilization mechanism 110 that is greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to retract tip stabilization mechanism 110 and fixation mechanism 130 relative to hypodermic tube 230. In one or more embodiments, advancing handle 210 towards eye 560 after a cannula contact 510 may actuate membrane removing tip distal end 241 into cannula distal end 571. Illustratively, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to actuate hypodermic tube distal end 231 into cannula distal end 571. In one or more embodiments, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to actuate membrane removing tip distal end 271 out from cannula proximal end 572. Illustratively, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to actuate hypodermic tube distal end 231 out from cannula proximal end 572. In one or more embodiments, advancing handle 210 towards eye 560 after a cannula contact 510 may be configured to actuate membrane removing tip 240 into inner eye 565.

FIG. 5D illustrates a complete membrane removing tip insertion 530. In one or more embodiments, advancing handle 210 towards eye 560 after a partial membrane removing tip insertion 520 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism distal end 111 towards handle distal end 211. In one or more embodiments, advancing handle 210 towards eye 560 after a partial membrane removing tip insertion 520 may be configured to apply a force to tip stabilization mechanism 110 that is greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, advancing handle 210 towards eye 560 after a partial membrane removing tip insertion 520 may be configured to retract tip stabilization mechanism 110 and fixation mechanism 130 relative to hypodermic tube 230. In one or more embodiments, advancing handle 210 towards eye 560 after a partial membrane removing tip insertion 520 may be configured to actuate membrane removing tip 240 within inner eye 565.

FIG. 5E illustrates a partial membrane removing tip extraction 540. Illustratively, a surgeon may guide a portion

of assembled instrument **300** out from cannula **570**, e.g., by retracting handle **210** away from eye **560** after a complete membrane removing tip insertion **530**. In one or more embodiments, retracting handle **210** away from eye **560** may be configured to reduce a force applied to tip stabilization mechanism **110**. Illustratively, reducing a force applied to tip stabilization mechanism **110** may be configured to fix tip stabilization mechanism **110** and fixation mechanism **130** in a position relative to hypodermic tube **230**. In one or more embodiments, a surgeon may use a distance between tip stabilization mechanism proximal end **112** and cannula distal end **571** as a depth gauge to evaluate a risk associated with a surgical procedure, e.g., during a difficult membrane removal, a surgeon may be concerned about damaging a patient's retinal tissue. Illustratively, a surgeon may use a distance between tip stabilization mechanism proximal end **112** and cannula distal end **571** as a depth gauge to compare a depth into the posterior segment of inner eye **565** of a particular portion of a surgical procedure with a depth into the posterior segment of inner eye **565** of a prior portion of a surgical procedure. For example, if a surgeon notices that a first attempt to raise a membrane from a retinal tissue causes damage to the retinal tissue, then the surgeon may note a depth of the first attempt to raise the membrane by observing a relative distance between tip stabilization mechanism proximal end **112** and cannula distal end **571**. Illustratively, the surgeon may make a second attempt to raise the membrane from the retinal tissue at a depth less than the depth of the first attempt to raise the membrane from the retinal tissue by preventing a contact between tip stabilization mechanism proximal end **112** and cannula distal end **571**.

FIG. 5F illustrates a complete membrane removing tip extraction **550**. Illustratively, a surgeon may guide a portion of assembled instrument **300** out from cannula **570**, e.g., by retracting handle **210** away from eye **560** after a partial membrane removing tip extraction **540**. In one or more embodiments, retracting handle **210** away from eye **560** after a partial membrane removing tip extraction **540** may be configured to extract membrane removing tip **240** from cannula **570**. Illustratively, after a complete membrane removing tip extraction **550**, tip stabilization mechanism **110** may be fixed in a position relative to hypodermic tube **230**, e.g., assembled instrument **300** may comprise an exposed membrane removing tip **400** after a complete membrane removing tip extraction **550**. In one or more embodiments, a surgeon may prepare assembled instrument **300** for a second cannula approach **500** after a complete membrane removing tip extraction **550** by grasping tip stabilization mechanism **110** and actuating tip stabilization mechanism **110** relative to hypodermic tube **230** towards membrane removing tip **240**. Illustratively, a surgeon may prepare assembled instrument **300** for a second cannula approach **500** after a complete membrane removing tip extraction **550** by actuating tip stabilization mechanism **110** over membrane removing tip **240**.

FIG. 6 is a schematic diagram illustrating an exploded view of an irrigating and aspirating instrument assembly **600**. In one or more embodiments, an irrigating and aspirating instrument assembly may comprise a tip stabilization mechanism **110**, a fixation mechanism **130**, a hypodermic tube **230**, a flow control mechanism **610**, a pressure vent tactile locater **620**, a fluid guide **630**, and an irrigating and aspirating tip **640**. Illustratively, flow control mechanism **610** may comprise a flow control mechanism distal end **611**, a flow control mechanism proximal end **612**, and a pressure vent **615**. In one or more embodiments, flow control mecha-

nism **610** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, fluid guide **630** may comprise a fluid guide distal end **631**, a fluid guide proximal end **632**, a distal locking lip **633**, a proximal locking lip **634**, and a locking depression **635**. In one or more embodiments, fluid guide **630** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, irrigating and aspirating tip **640** may comprise an irrigating and aspirating tip distal end **641** and an irrigating and aspirating tip proximal end **642**. In one or more embodiments, irrigating and aspirating tip **640** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials. Illustratively, irrigating and aspirating tip **640** may be manufactured from silicone, e.g., biocompatible silicone. In one or more embodiments, irrigating and aspirating tip **640** may comprise an inner lumen for irrigation of fluids. Illustratively, irrigating and aspirating tip **640** may comprise an inner lumen for aspiration of fluids. In one or more embodiments, irrigating and aspirating tip **640** may comprise an inner lumen for irrigation of fluids and for aspiration of fluids.

FIGS. 7A and 7B are schematic diagrams illustrating an assembled irrigating and aspirating instrument **700**. FIG. 7A illustrates a top view of an assembled irrigating and aspirating instrument **700**. In one or more embodiments, fixation mechanism **130** may be disposed over a portion of tip stabilization mechanism **110**, e.g., fixation mechanism **130** may be disposed between tip stabilization mechanism distal end **111** and tip stabilization mechanism proximal end **112**. Illustratively, a portion of fixation mechanism **130** may be disposed within a portion of tip stabilization mechanism **110**, e.g., a portion of fixation mechanism **130** may be disposed within fixation mechanism channel **115**. In one or more embodiments, a portion of fixation mechanism **130** may be disposed within inner bore **120**. Illustratively, a portion of fixation mechanism **130** may be fixed to a portion of tip stabilization mechanism **110**, e.g., fixation mechanism inner diameter **136** may be fixed to a portion of tip stabilization mechanism **110**. In one or more embodiments, a portion of fixation mechanism **130** may be fixed to a portion of tip stabilization mechanism **110** by a force of friction or by any suitable fixation means, e.g., a portion of fixation mechanism **130** may be fixed to a portion of tip stabilization mechanism **110** by an adhesive, a weld, etc.

Illustratively, tip stabilization mechanism **110** may be disposed over a portion of hypodermic tube **230**, e.g., tip stabilization mechanism distal end **111** may be disposed over hypodermic tube distal end **231**. In one or more embodiments, a portion of hypodermic tube **230** may be disposed within inner bore **120**, e.g., hypodermic tube distal end **231** may be disposed within inner bore **120**. Illustratively, hypodermic tube **230** may be disposed within fixation mechanism **130**, e.g., hypodermic tube **230** may be disposed within fixation mechanism inner diameter **136**. In one or more embodiments, hypodermic tube **230** may not be disposed within fixation mechanism **130**, e.g., a portion of hypodermic tube **230** may be disposed adjacent to a portion of fixation mechanism **130**. Illustratively, fixation mechanism **130** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**, e.g., a static friction force between fixation mechanism **130** and hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**. In one or more embodi-

ments, a contact between a portion of fixation mechanism **130** and a portion of hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**. Illustratively, fixation mechanism **130** may be fixed to both tip stabilization mechanism **110** and hypodermic tube **230** wherein a static friction force between fixation mechanism **130** and hypodermic tube **230** may be configured to temporarily fix tip stabilization mechanism **110** in a position relative to hypodermic tube **230**.

Illustratively, a portion of irrigating and aspirating tip **640** may be disposed within hypodermic tube **230**, e.g., irrigating and aspirating tip proximal end **642** may be disposed within hypodermic tube distal end **231**. In one or more embodiments, a portion of irrigating and aspirating tip **640** may be disposed within hypodermic tube **230** wherein a portion of irrigating and aspirating tip **640** extends from a portion of hypodermic tube **230**, e.g., a portion of irrigating and aspirating tip **640** may be disposed within hypodermic tube **230** wherein irrigating and aspirating tip distal end **641** extends from hypodermic tube distal end **231**. Illustratively, a portion of irrigating and aspirating tip **640** may be fixed within hypodermic tube **230**, e.g., irrigating and aspirating tip proximal end **642** may be fixed within hypodermic tube **230**. In one or more embodiments, a portion of irrigating and aspirating tip **640** may be fixed within hypodermic tube **230** by an adhesive or any suitable fixation means, e.g., a portion of irrigating and aspirating tip **640** may be fixed within hypodermic tube **230** by a friction fit, a weld, etc.

Illustratively, pressure vent tactile locater **620** may be disposed over a portion of flow control mechanism **610**, e.g., pressure vent tactile locater **620** may be disposed over pressure vent **615**. In one or more embodiments, pressure vent tactile locater **620** may be fixed to a portion of flow control mechanism **610** by an adhesive or any suitable fixation means. Illustratively, pressure vent tactile locater **620** may be configured to allow a surgeon to tactilely identify a location of pressure vent **615** where visual identification of a location of pressure vent **615** is impractical, e.g., in a dark operating room. In one or more embodiments, pressure vent tactile locater **620** may be configured to indicate one or more properties of assembled irrigating and aspirating instrument **700** to a surgeon, a nurse, or a surgical technician, e.g., pressure vent tactile locater **620** may comprise a color or a marking configured to visually indicate a minimum cannula gauge size that hypodermic tube **230** may ingress to perform a surgical procedure. Illustratively, pressure vent tactile locater **620** may be manufactured from any suitable material, e.g., polymers, metals, metal alloys, etc., or from any combination of suitable materials.

FIG. 7B illustrates a cross-sectional view of an assembled irrigating and aspirating instrument **700**. Illustratively, assembled irrigating and aspirating instrument **700** may comprise a fluid chamber **710**, a fluid conduit **720**, an irrigation taper **730**, and a hypodermic tube housing **740**. In one or more embodiments, a portion of fluid guide **630** may be disposed within flow control mechanism **610**, e.g., fluid guide proximal end **632** may be disposed within flow control mechanism **610**. Illustratively, a portion of fluid guide **630** may be fixed within a flow control mechanism **610** by any suitable fixation mechanism, e.g., a portion of fluid guide **630** may be fixed within flow control mechanism **610** by an adhesive, a friction fit, a locking fit, a weld, etc. In one or more embodiments a portion of hypodermic tube **230** may be disposed within a portion of fluid guide **630**, e.g., hypodermic tube proximal end **232** may be disposed within hypodermic tube housing **740**. Illustratively, a portion of

hypodermic tube **230** may be fixed within hypodermic tube housing **740** by any suitable fixation mechanism, e.g., a portion of hypodermic tube **230** may be fixed within hypodermic tube housing **740** by a press-fit, an adhesive, a weld, etc.

Illustratively, irrigating and aspirating tip **640** may be disposed within hypodermic tube **230** and tip stabilization mechanism **110**, e.g., irrigating and aspirating tip distal end **641** may be disposed between tip stabilization mechanism distal end **111** and tip stabilization mechanism proximal end **112**. In one or more embodiments, irrigating and aspirating tip **640** may be disposed within hypodermic tube **230** and tip stabilization mechanism **110** wherein irrigating and aspirating tip proximal end **642** may be disposed between hypodermic tube distal end **231** and hypodermic tube proximal end **232**. Illustratively, hypodermic tube **230** may be disposed within tip stabilization mechanism **110** wherein hypodermic tube distal end **231** is disposed between tip stabilization mechanism distal end **111** and tip stabilization mechanism proximal end **112**.

FIG. 8 is a schematic diagram illustrating an exposed irrigating and aspirating tip **800**. In one or more embodiments, irrigating and aspirating tip **640** may be colored black, e.g., to enhance visualization of irrigating and aspirating tip **640**. For example, irrigating and aspirating tip **640** may be manufactured from black silicone to enhance visualization of irrigating and aspirating tip **640**. Illustratively, irrigating and aspirating tip **640** may be colored any color not contained within Johannes Itten's color wheel to enhance visualization of irrigating and aspirating tip **640**, e.g., irrigating and aspirating tip **640** may be colored black or grey to enhance visualization of irrigating and aspirating tip **640**. In one or more embodiments, irrigating and aspirating tip **640** may be configured to irrigate fluid into inner eye **565**. Illustratively, irrigating and aspirating tip **640** may be configured to aspirate fluid out from inner eye **565**. In one or more embodiments, irrigating and aspirating tip **640** may be configured to both irrigate fluid into inner eye **565** and to aspirate fluid out from inner eye **565**.

FIGS. 9A, 9B, 9C, 9D, 9E, and 9F are schematic diagrams illustrating a portion of a surgical procedure. FIG. 9A illustrates a cannula approach **900**. Illustratively, a surgeon may maneuver an assembled irrigating and aspirating instrument **700** towards an eye **560** to perform a portion of a surgical procedure, e.g., a surgeon may maneuver an assembled irrigating and aspirating instrument **700** towards a cannula **570** to perform a portion of a surgical procedure. In one or more embodiments, cannula **570** may comprise a cannula distal end **571** and a cannula proximal end **572**. Illustratively, cannula **570** may be disposed within an incision in eye **560**, e.g., cannula **570** may be disposed within an incision in eye **560** wherein cannula distal end **571** extends out from an outer surface of eye **560** and cannula proximal end **572** is disposed within an inner eye **565**. In one or more embodiments, a cannula approach **900** may comprise an attempt to guide tip stabilization mechanism proximal end **112** towards cannula distal end **571**.

FIG. 9B illustrates a cannula contact **910**. Illustratively, a surgeon may approach cannula **570** with assembled irrigating and aspirating instrument **700** until a portion of assembled irrigating and aspirating instrument **700** contacts a portion of cannula **570**, e.g., a surgeon may approach cannula **570** with assembled irrigating and aspirating instrument **700** until tip stabilization mechanism proximal end **112** contacts cannula distal end **571**. In one or more embodiments, a cannula contact **510** may comprise a contact between tip stabilization mechanism proximal end **112** and

cannula distal end 571. Illustratively, a cannula contact 510 may be configured to apply a force to fixation mechanism 130, e.g., a cannula contact 510 may apply a normal force to tip stabilization mechanism 110. In one or more embodiments, an application of a normal force to tip stabilization mechanism 110 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, an application of a normal force to tip stabilization mechanism 110 may not be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be less than a static friction force between fixation mechanism 130 and hypodermic tube 230.

FIG. 9C illustrates a partial irrigating and aspirating tip insertion 920. Illustratively, a surgeon may guide a portion of assembled irrigating and aspirating instrument 700 into cannula 570, e.g., by advancing flow control mechanism 610 towards eye 560 after a cannula contact 510. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism distal end 111 towards handle distal end 211. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to apply a force to tip stabilization mechanism 110 that is greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to retract tip stabilization mechanism 110 and fixation mechanism 130 relative to hypodermic tube 230. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may actuate irrigating and aspirating tip distal end 641 into cannula distal end 571. Illustratively, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to actuate hypodermic tube distal end 231 into cannula distal end 571. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to actuate irrigating and aspirating tip distal end 641 out from cannula proximal end 572. Illustratively, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to actuate hypodermic tube distal end 231 out from cannula proximal end 572. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a cannula contact 510 may be configured to actuate irrigating and aspirating tip 640 into inner eye 565.

FIG. 9D illustrates a complete irrigating and aspirating tip insertion 930. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a partial irrigating and aspirating tip insertion 920 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism distal end 111 towards

fluid guide distal end 631. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a partial irrigating and aspirating tip insertion 920 may be configured to apply a force to tip stabilization mechanism 110 that is greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, advancing flow control mechanism 610 towards eye 560 after a partial irrigating and aspirating tip insertion 920 may be configured to retract tip stabilization mechanism 110 and fixation mechanism 130 relative to hypodermic tube 230. In one or more embodiments, advancing flow control mechanism 610 towards eye 560 after a partial irrigating and aspirating tip insertion 920 may be configured to actuate irrigating and aspirating tip 640 within inner eye 565.

FIG. 9E illustrates a partial irrigating and aspirating tip extraction 940. Illustratively, a surgeon may guide a portion of assembled irrigating and aspirating instrument 700 out from cannula 570, e.g., by retracting flow control mechanism 610 away from eye 560 after a complete irrigating and aspirating tip insertion 930. In one or more embodiments, retracting flow control mechanism 610 away from eye 560 may be configured to reduce a force applied to tip stabilization mechanism 110. Illustratively, reducing a force applied to tip stabilization mechanism 110 may be configured to fix tip stabilization mechanism 110 and fixation mechanism 130 in a position relative to hypodermic tube 230. In one or more embodiments, a surgeon may use a distance between tip stabilization mechanism proximal end 112 and cannula distal end 571 as a depth gauge to evaluate a risk associated with a surgical procedure, e.g., during a difficult membrane removal, a surgeon may be concerned about damaging a patient's retinal tissue. Illustratively, a surgeon may use a distance between tip stabilization mechanism proximal end 112 and cannula distal end 571 as a depth gauge to compare a depth into the posterior segment of inner eye 565 of a particular portion of a surgical procedure with a depth into the posterior segment of inner eye 565 of a prior portion of a surgical procedure.

FIG. 9F illustrates a complete irrigating and aspirating tip extraction 950. Illustratively, a surgeon may guide a portion of assembled irrigating and aspirating instrument 700 out from cannula 570, e.g., by retracting flow control mechanism 610 away from eye 560 after a partial irrigating and aspirating tip extraction 940. In one or more embodiments, retracting flow control mechanism 610 away from eye 560 after a partial irrigating and aspirating tip extraction 940 may be configured to extract irrigating and aspirating tip 610 from cannula 570. Illustratively, after a complete irrigating and aspirating tip extraction 950, tip stabilization mechanism 110 may be fixed in a position relative to hypodermic tube 230, e.g., assembled irrigating and aspirating instrument 700 may comprise an exposed irrigating and aspirating tip 800 after a complete irrigating and aspirating tip extraction 950. In one or more embodiments, a surgeon may prepare assembled irrigating and aspirating instrument 700 for a second cannula approach 900 after a complete irrigating and aspirating tip extraction 950 by grasping tip stabilization mechanism 110 and actuating tip stabilization mechanism 110 relative to hypodermic tube 230 towards irrigating and aspirating tip 640. Illustratively, a surgeon may prepare assembled irrigating and aspirating instrument 700 for a second cannula approach 900 after a complete irrigating and aspirating tip extraction 950 by actuating tip stabilization mechanism 110 over irrigating and aspirating tip 640.

FIGS. 10A, 10B, 10C, 10D, and 10E are schematic diagrams illustrating a portion of a surgical procedure. FIG.

10A illustrates a valved cannula approach 1000. Illustratively, a valved cannula 1050 may comprise an inner lumen 1055, a first valve hinge 1065, a second valve hinge 1065, and a valve hinge interface 1060. In one or more embodiments, first valve hinge 1065 may comprise a first valve hinge superior surface 1066 and second valve hinge 1065 may comprise a second valve hinge superior surface 1066. Illustratively, valved cannula 1050 may be configured to maintain an intraocular pressure, e.g., valve hinge interface 1060 may be configured to maintain an intraocular pressure during a surgical procedure. In one or more embodiments, a surgeon may guide tip stabilization mechanism 110 towards valved cannula 1050 to perform a portion of a surgical procedure, e.g., a surgeon may guide tip stabilization mechanism proximal end 112 towards valve hinge superior surface 1066 to perform a portion of a surgical procedure. Illustratively, as a surgeon guides tip stabilization mechanism proximal end 112 towards valve hinge superior surface 1066, hypodermic tube distal end 231 may be disposed within tip stabilization mechanism inner bore 120. In one or more embodiments, as a surgeon guides tip stabilization mechanism proximal end 112 towards valve hinge superior surface 1066, membrane removing tip distal end 241 may be disposed within tip stabilization mechanism inner bore 120.

FIG. 10B illustrates a valved cannula contact 1010. In one or more embodiments, a valved cannula contact 1010 may comprise a contact between a portion of tip stabilization mechanism 110 and a portion of valved cannula 1050, e.g., a valved cannula contact 1010 may comprise a contact between tip stabilization mechanism proximal end 112 and valve hinge superior surface 1066. Illustratively, a valved cannula contact 1010 may be configured to apply a force to fixation mechanism 130, e.g., a valved cannula contact 1010 may apply a normal force to tip stabilization mechanism 110. In one or more embodiments, an application of a normal force to tip stabilization mechanism 110 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, an application of a normal force to tip stabilization mechanism 110 may not be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be less than a static friction force between fixation mechanism 130 and hypodermic tube 230.

FIG. 10C illustrates a membrane removing tip stabilization 1020. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after a valved cannula contact 1010, e.g., a surgeon may actuate hypodermic tube distal end 231 towards inner lumen 1055 after a valved cannula contact 1010. In one or more embodiments, actuating hypodermic tube 230 towards inner lumen 1055 after a valved cannula contact 1010 may be configured to apply a force to tip stabilization mechanism 110, e.g., actuating hypodermic tube distal end 231 towards inner lumen 1055 after a valved cannula contact 1010 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism 110 may be configured to actuate tip stabilization mechanism 110 relative to housing tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 that is greater than a force of static friction between fixation mechanism 130 and hypodermic tube 230 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230. Illustratively, an application of a force to tip stabilization mechanism proximal end

112 may be configured to actuate tip stabilization mechanism distal end 111 towards hypodermic tube proximal end 232. In one or more embodiments, an actuation of tip stabilization mechanism 110 relative to housing tube 230 may be configured to expose a portion of membrane removing tip 240, e.g., an actuation of tip stabilization mechanism distal end 111 towards hypodermic tube proximal end 232 may be configured to expose membrane removing tip distal end 241. Illustratively, an exposure of a portion of membrane removing tip 240 may be configured to facilitate a contact between membrane removing tip 240 and valve hinge superior surface 1066, e.g., an exposure of a portion of membrane removing tip 240 may be configured to facilitate a contact between membrane removing tip distal end 241 and valve hinge superior surface 1066. In one or more embodiments, as a surgeon actuates hypodermic tube distal end 231 towards inner lumen 1055, a contact between membrane removing tip 240 and valve hinge superior surface 1066 may be configured to cause membrane removing tip 240 to deform, e.g., a contact between membrane removing tip 240 and valve hinge superior surface 1066 may be configured to cause membrane removing tip 240 to bend or flex. Illustratively, a contact between membrane removing tip 240 and valve hinge superior surface 1066 may be configured to cause an elastic deformation of membrane removing tip 240. In one or more embodiments, a contact between membrane removing tip 240 and valve hinge superior surface 1066 may be configured to cause a plastic deformation of membrane removing tip 240.

Illustratively, tip stabilization mechanism 110 may be configured to facilitate a membrane removing tip deformation prevention 1070, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent membrane removing tip 240 from deforming. In one or more embodiments, tip stabilization mechanism 110 may be configured to prevent a portion of membrane removing tip 240 from deforming beyond a line tangent to a wall of inner lumen 1055, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent a portion of membrane removing tip 240 from deforming beyond a line tangent to a wall of inner lumen 1055. For example, tip stabilization mechanism 110 may be configured to prevent a portion of membrane removing tip 240 from bending or flexing beyond a line tangent to a wall of inner lumen 1055. Illustratively, tip stabilization mechanism 110 may be configured to prevent a portion of membrane removing tip 240 from deforming beyond a line tangent to an outer diameter of valved cannula 1050, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent a portion of membrane removing tip 240 from deforming beyond a line tangent to an outer diameter of valved cannula 1050. For example, tip stabilization mechanism 110 may be configured to prevent a portion of membrane removing tip 240 from bending or flexing beyond a line tangent to an outer diameter of valved cannula 1050. In one or more embodiments, a membrane removing tip deformation prevention 1070 may be configured to actuate valve hinge 1065, e.g., a membrane removing tip deformation prevention 1070 may be configured to separate a first valve hinge 1065 and a second valve hinge 1065 at valve hinge interface 1060. Illustratively, a membrane removing tip deformation prevention 1070 may be configured to guide a portion of membrane removing tip 240 into inner lumen 1055, e.g., a membrane removing tip deformation prevention 1070 may be configured to guide membrane removing tip distal end 241 into inner lumen 1055. In one or more embodiments, membrane removing tip 240 may have a first stiffness, valve hinge 1065 may have a

second stiffness, and tip stabilization mechanism 110 may have a third stiffness wherein the second stiffness is greater than the first stiffness and the third stiffness is greater than the second stiffness. Illustratively, the second stiffness of valve hinge 1065 may be configured to cause a deformation of membrane removing tip 240 and the third stiffness of tip stabilization mechanism 110 may be configured to guide a portion of membrane removing tip 240 into inner lumen 1055.

FIG. 10D illustrates a partial valved cannula ingress 1030. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after a membrane removing tip stabilization 1020, e.g., a surgeon may actuate hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020. In one or more embodiments, an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020 may be configured to actuate a portion of membrane removing tip 240 into valved cannula 1050, e.g., an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020 may be configured to actuate abrasive surface 410 into inner lumen 1055. In one or more embodiments, an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020 may be configured to increase a separation of a first valve hinge 1065 and a second valve hinge 1065. Illustratively, an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020 may be configured to cause membrane removing tip 240 to return to its original geometry, e.g., an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after a membrane removing tip stabilization 1020 may be configured to cause membrane removing tip 240 to return to its pre-deformation geometry.

FIG. 10E illustrates a complete valved cannula ingress 1040. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after a partial valved cannula ingress 1030, e.g., a surgeon may actuate hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1030. In one or more embodiments, an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1030 may be configured to fully separate a first valve hinge 1065 and a second valve hinge 1065. Illustratively, an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1030 may be configured to cause a portion of hypodermic tube 230 to contact a portion of valve hinge superior surface 1066, e.g., an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1030 may be configured to cause an outer portion of hypodermic tube 230 to contact a portion of valve hinge superior surface 1066.

FIGS. 11A, 11B, 11C, 11D, and 11E are schematic diagrams illustrating a portion of a surgical procedure. FIG. 11A illustrates a valved cannula approach 1100. In one or more embodiments, a surgeon may guide tip stabilization mechanism 110 towards valved cannula 1050 to perform a portion of a surgical procedure, e.g., a surgeon may guide tip stabilization mechanism proximal end 112 towards valve hinge superior surface 1066 to perform a portion of a surgical procedure. Illustratively, as a surgeon guides tip stabilization mechanism proximal end 112 towards valve

hinge superior surface 1066, hypodermic tube distal end 231 may be disposed within tip stabilization mechanism inner bore 120. In one or more embodiments, as a surgeon guides tip stabilization mechanism proximal end 112 towards valve hinge superior surface 1066, irrigating and aspirating tip distal end 641 may be disposed within tip stabilization mechanism inner bore 120.

FIG. 11B illustrates a valved cannula contact 1110. In one or more embodiments, a valved cannula contact 1110 may comprise a contact between a portion of tip stabilization mechanism 110 and a portion of valved cannula 1050, e.g., a valved cannula contact 1110 may comprise a contact between tip stabilization mechanism proximal end 112 and valve hinge superior surface 1066. Illustratively, a valved cannula contact 1110 may be configured to apply a force to fixation mechanism 130, e.g., a valved cannula contact 1110 may apply a normal force to tip stabilization mechanism 110. In one or more embodiments, an application of a normal force to tip stabilization mechanism 110 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be greater than a static friction force between fixation mechanism 130 and hypodermic tube 230. Illustratively, an application of a normal force to tip stabilization mechanism 110 may not be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230, e.g., a normal force applied to tip stabilization mechanism 110 may be less than a static friction force between fixation mechanism 130 and hypodermic tube 230.

FIG. 11C illustrates an irrigating and aspirating tip stabilization 1120. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after a valved cannula contact 1110, e.g., a surgeon may actuate hypodermic tube distal end 231 towards inner lumen 1055 after a valved cannula contact 1110. In one or more embodiments, actuating hypodermic tube 230 towards inner lumen 1055 after a valved cannula contact 1110 may be configured to apply a force to tip stabilization mechanism 110, e.g., actuating hypodermic tube distal end 231 towards inner lumen 1055 after a valved cannula contact 1110 may be configured to apply a force to tip stabilization mechanism proximal end 112. Illustratively, an application of a force to tip stabilization mechanism 110 may be configured to actuate tip stabilization mechanism 110 relative to housing tube 230, e.g., an application of a force to tip stabilization mechanism proximal end 112 that is greater than a force of static friction between fixation mechanism 130 and hypodermic tube 230 may be configured to retract tip stabilization mechanism 110 relative to hypodermic tube 230. Illustratively, an application of a force to tip stabilization mechanism proximal end 112 may be configured to actuate tip stabilization mechanism distal end 111 towards hypodermic tube proximal end 232. In one or more embodiments, an actuation of tip stabilization mechanism 110 relative to housing tube 230 may be configured to expose a portion of irrigating and aspirating tip 640, e.g., an actuation of tip stabilization mechanism distal end 111 towards hypodermic tube proximal end 232 may be configured to expose irrigating and aspirating tip distal end 641. Illustratively, an exposure of a portion of irrigating and aspirating tip 640 may be configured to facilitate a contact between irrigating and aspirating tip 640 and valve hinge superior surface 1066, e.g., an exposure of a portion of irrigating and aspirating tip 640 may be configured to facilitate a contact between irrigating and aspirating tip distal end 641 and valve hinge superior surface 1066. In one or more embodiments, as a surgeon actuates hypodermic tube distal end 231 towards

inner lumen 1055, a contact between irrigating and aspirating tip 640 and valve hinge superior surface 1066 may be configured to cause irrigating and aspirating tip 640 to deform, e.g., a contact between irrigating and aspirating tip 640 and valve hinge superior surface 1066 may be configured to cause irrigating and aspirating tip 640 to bend or flex. Illustratively, a contact between irrigating and aspirating tip 640 and valve hinge superior surface 1066 may be configured to cause an elastic deformation of irrigating and aspirating tip 640. In one or more embodiments, a contact between irrigating and aspirating tip 640 and valve hinge superior surface 1066 may be configured to cause a plastic deformation of irrigating and aspirating tip 640.

Illustratively, tip stabilization mechanism 110 may be configured to facilitate an irrigating and aspirating tip deformation prevention 1170, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent irrigating and aspirating tip 640 from deforming. In one or more embodiments, tip stabilization mechanism 110 may be configured to prevent a portion of irrigating and aspirating tip 640 from deforming beyond a line tangent to a wall of inner lumen 1055, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent a portion of irrigating and aspirating tip 640 from deforming beyond a line tangent to a wall of inner lumen 1055. For example, tip stabilization mechanism 110 may be configured to prevent a portion of irrigating and aspirating tip 640 from bending or flexing beyond a line tangent to a wall of inner lumen 1055. Illustratively, tip stabilization mechanism 110 may be configured to prevent a portion of irrigating and aspirating tip 640 from deforming beyond a line tangent to an outer diameter of valved cannula 1050, e.g., tip stabilization mechanism inner diameter 114 may be configured to prevent a portion of irrigating and aspirating tip 640 from deforming beyond a line tangent to an outer diameter of valved cannula 1050. For example, tip stabilization mechanism 110 may be configured to prevent a portion of irrigating and aspirating tip 640 from bending or flexing beyond a line tangent to an outer diameter of valved cannula 1050. In one or more embodiments, an irrigating and aspirating tip deformation prevention 1170 may be configured to actuate valve hinge 1065, e.g., an irrigating and aspirating tip deformation prevention 1170 may be configured to separate a first valve hinge 1065 and a second valve hinge 1065 at valve hinge interface 1060. Illustratively, an irrigating and aspirating tip deformation prevention 1170 may be configured to guide a portion of irrigating and aspirating tip 640 into inner lumen 1055, e.g., a membrane removing tip deformation prevention 1070 may be configured to guide irrigating and aspirating tip distal end 641 into inner lumen 1055. In one or more embodiments, irrigating and aspirating tip 640 may have a first stiffness, valve hinge 1065 may have a second stiffness, and tip stabilization mechanism 110 may have a third stiffness wherein the second stiffness is greater than the first stiffness and the third stiffness is greater than the second stiffness. Illustratively, the second stiffness of valve hinge 1065 may be configured to cause a deformation of irrigating and aspirating tip 640 and the third stiffness of tip stabilization mechanism 110 may be configured to guide a portion of irrigating and aspirating tip 640 into inner lumen 1055.

FIG. 11D illustrates a partial valved cannula ingress 1130. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120, e.g., a surgeon may actuate hypodermic tube distal end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120. In one or more embodiments, an actuation of hypodermic tube distal

end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120 may be configured to actuate a portion of irrigating and aspirating tip 640 into valved cannula 1050, e.g., an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120 may be configured to actuate irrigating and aspirating tip distal end 641 into inner lumen 1055. Illustratively, an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120 may be configured to increase a separation of a first valve hinge 1065 and a second valve hinge 1065. Illustratively, an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120 may be configured to cause irrigating and aspirating tip 640 to return to its original geometry, e.g., an actuation of hypodermic tube distal end 231 towards inner lumen 1055 after an irrigating and aspirating tip stabilization 1120 may be configured to cause irrigating and aspirating tip 640 to return to its pre-deformation geometry.

FIG. 11E illustrates a complete valved cannula ingress 1140. Illustratively, a surgeon may actuate hypodermic tube 230 towards inner lumen 1055 after a partial valved cannula ingress 1130, e.g., a surgeon may actuate hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1130. In one or more embodiments, an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1130 may be configured to fully separate a first valve hinge 1065 and a second valve hinge 1065. Illustratively, an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1130 may be configured to cause a portion of hypodermic tube 230 to contact a portion of valve hinge superior surface 1066, e.g., an actuation of hypodermic tube distal end 231 into inner lumen 1055 after a partial valved cannula ingress 1130 may be configured to cause an outer portion of hypodermic tube 230 to contact a portion of valve hinge superior surface 1066.

The foregoing description has been directed to particular embodiments of this invention. It will be apparent; however, that other variations and modifications may be made to the described embodiments, with the attainment of some or all of their advantages. Specifically, it should be noted that the principles of the present invention may be implemented in any system. Furthermore, while this description has been written in terms of a cannula ingress system, the teachings of the present invention are equally suitable to any systems where the functionality may be employed. Therefore, it is the object of the appended claims to cover all such variations and modifications as come within the true spirit and scope of the invention.

What is claimed is:

1. A system comprising:

- a hypodermic tube having a hypodermic tube distal end and a hypodermic tube proximal end, the hypodermic tube having dimensions configured for performing ophthalmic surgery;
- a tip having a tip distal end and a tip proximal end wherein the tip proximal end is disposed within the hypodermic tube and the tip distal end extends out from the hypodermic tube distal end;
- a tip stabilization mechanism having a tip stabilization mechanism distal end and a tip stabilization mechanism proximal end; and
- a fixation mechanism disposed over a portion of the tip stabilization mechanism wherein a portion of the fixation mechanism is disposed in a fixation mechanism

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channel of the tip stabilization mechanism wherein the tip distal end and the hypodermic tube distal end are disposed within an inner bore of the tip stabilization mechanism and the fixation mechanism is configured to temporarily fix the tip stabilization mechanism in a position relative to the hypodermic tube wherein the tip stabilization mechanism has an outer diameter that is less than an outer diameter of the fixation mechanism.

2. The system of claim 1 wherein the fixation mechanism contacts a portion of the hypodermic needle.

3. The system of claim 2 wherein a static friction force is configured to temporarily fix the tip stabilization mechanism in the position relative to the hypodermic tube.

4. The system of claim 1 wherein the tip stabilization mechanism is configured to prevent the tip from deforming beyond a line tangent to an outer diameter of a valved cannula.

5. The system of claim 4 wherein the tip stabilization mechanism is configured to actuate a first valve hinge of the valved cannula.

6. The system of claim 5 wherein the tip stabilization mechanism is configured to separate the first valve hinge and a second valve hinge at a valve hinge interface.

7. The system of claim 6 wherein the tip stabilization mechanism is configured to guide the tip distal end into an inner lumen of the valved cannula.

8. The system of claim 1 wherein the tip stabilization mechanism is configured to prevent the tip from deforming beyond a line tangent to a wall of an inner lumen of a cannula.

9. The system of claim 1 wherein the tip stabilization mechanism is configured to prevent the tip from deforming after a contact between the tip distal end and a valve hinge superior surface.

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10. The system of claim 1 further comprising: an abrasive surface of the tip configured to grasp a portion of a membrane.

11. The system of claim 10 further comprising: biocompatible particles of the abrasive surface.

12. The system of claim 11 wherein the tip is colored black to enhance visualization of the biocompatible particles of the abrasive surface.

13. The system of claim 1 wherein the tip is colored black to enhance visualization of the tip.

14. The system of claim 13 wherein the tip is manufactured from black silicone.

15. The system of claim 1 wherein the tip is colored grey to enhance visualization of the tip.

16. The system of claim 1 wherein the tip is colored any color not contained within Johannes Itten's color wheel to enhance visualization of the tip.

17. The system of claim 1 further comprising: an inner lumen of the tip configured for irrigation of fluids.

18. The system of claim 17 wherein the inner lumen of the tip is configured for aspiration of fluids.

19. The system of claim 1 wherein an application of a force to the tip stabilization mechanism proximal end is configured to actuate the tip stabilization mechanism relative to the hypodermic tube.

20. The system of claim 19 wherein the application of the force to the tip stabilization mechanism proximal end is configured to expose the tip distal end.

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